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202 CGAGAAACCTGCGGATCGGCTCGCTAC 231

seq_name: gb_pr2:HSHLABB1

seq_documentation_block:
LOCUS      HSHLABB1      250 bp      DNA      PRI      25-MAR-1997
DEFINITION Human cell line THAI DCH010 MHC class I HLA-B gene (allele
HLA-B*51V), exon 2.
ACCESSION  U90611
VERSION     U90611.1 GI:1905865
KEYWORDS
SEGMENT    1 of 2
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 250)
AUTHORS    Chandanayingyong,D., Sirikong,M., Longta,K., Srinak,D.,
Siriboonrit,U., Rungroung,E. and Bejchandra,S.
TITLE      B*51V alleles
JOURNAL    Unpublished
REFERENCE  2 (bases 1 to 250)
AUTHORS    Chandanayingyong,D., Sirikong,M., Longta,K., Srinak,D.,
Siriboonrit,U., Rungroung,E. and Bejchandra,S.
TITLE      Direct Submission
JOURNAL    Submitted (25-FEB-1997) Transfusion Medicine, Faculty of Medicine,
Siriraj Hospital, Mahidol University, Prannok Road, Bangkok 10700,
Thailand
FEATURES   Location/Qualifiers
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Percent Similarity: 100.000 Percent Identity: 90.000

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Align seg 1/1 to: HSHLABB1 from: 1 to: 250

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seq_name: gb_pr2:HSHLABB1

seq_documentation_block:
LOCUS      HSHLABB1      250 bp      DNA      PRI      25-MAR-1997
DEFINITION Human cell line THAI DCH010 MHC class I HLA-B gene (allele
HLA-B*51V), exon 2.
ACCESSION  U90611
VERSION     U90611.1 GI:1905865
KEYWORDS
SEGMENT    1 of 2
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 250)
AUTHORS    Chandanayingyong,D., Sirikong,M., Longta,K., Srinak,D.,
Siriboonrit,U., Rungroung,E. and Bejchandra,S.
TITLE      B*51V alleles
JOURNAL    Unpublished
REFERENCE  2 (bases 1 to 250)
AUTHORS    Chandanayingyong,D., Sirikong,M., Longta,K., Srinak,D.,
Siriboonrit,U., Rungroung,E. and Bejchandra,S.
TITLE      Direct Submission
JOURNAL    Submitted (25-FEB-1997) Transfusion Medicine, Faculty of Medicine,
Siriraj Hospital, Mahidol University, Prannok Road, Bangkok 10700,
Thailand
FEATURES   Location/Qualifiers
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BASE COUNT  56 a 82 c 80 g 32 t
ORIGIN

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DEFINITION Human cell line THAI DCH028 MHC class I HLA-B gene (allele
HLA-B*51V), exon 2.
ACCESSION  U90613
VERSION     U90613.1 GI:1906033
KEYWORDS
SEGMENT    1 of 2
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 250)
AUTHORS    Chandanayingyong,D., Sirikong,M., Longta,K., Srinak,D.,
Siriboonrit,U., Rungroung,E. and Bejchandra,S.
TITLE      B*51V alleles
JOURNAL    Unpublished
REFERENCE  2 (bases 1 to 250)
AUTHORS    Chandanayingyong,D., Sirikong,M., Longta,K., Srinak,D.,
Siriboonrit,U., Rungroung,E. and Bejchandra,S.
TITLE      Direct Submission
JOURNAL    Submitted (25-FEB-1997) Transfusion Medicine, Faculty of Medicine,
Siriraj Hospital, Mahidol University, Prannok Road, Bangkok 10700,
Thailand
FEATURES   Location/Qualifiers
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Align seg 1/1 to: HSHLABG1 from: 1 to: 250

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seq_name: gb_pr2:HSHLABG1

seq_documentation_block:
LOCUS      HSHLABG1      250 bp      DNA      PRI      25-MAR-1997
DEFINITION Human cell line THAI DCH011 MHC class I HLA-B gene (allele
HLA-B*51V), exon 2.
ACCESSION  U90615

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KEYWORDS
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
            Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 259)
AUTHORS     Grimsley, C., Mather, K.A. and Ober, C.
TITLE       HLA-H: a pseudogene with increased variation due to balancing
            selection at neighboring loci
JOURNAL     Mol. Biol. Evol. 15 (12), 1581-1588 (1998)
MEDLINE     99083426
REFERENCE   2 (bases 1 to 259)
AUTHORS     Grimsley, C., Mather, K.A. and Ober, C.
TITLE       Direct Submission
JOURNAL     Submitted (03-SEP-1997) Fred Hutchinson Cancer Research Center,
            1100 Fairview Ave. N., M374, Seattle, WA 98109, USA
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  DEFINITION Sequence 67 from patent US 5451512.
  ACCESSION  II4590
  VERSION     II4590.1 GI:997073
  KEYWORDS   Unknown.
  SOURCE      Unknown.
  ORGANISM    Unknown.
  UNCLASSIFIED
  REFERENCE   1 (bases 1 to 270)
  AUTHORS     Apple, R.J., Bugawan, T.L. and Erlich, H.A.
  TITLE       Methods and reagents for HLA class I A locus DNA typing
  JOURNAL     Patent: US 5451512-A 67 19-SEP-1995;
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BASE COUNT  55 a   84 c   95 g   36 t
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  Ratio:     4.400     Gaps:     0
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  ORGANISM    Unknown.
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  REFERENCE   1 (bases 1 to 270)
  AUTHORS     Apple, R.J., Bugawan, T.L. and Erlich, H.A.
  TITLE       Methods and reagents for HLA class I A locus DNA typing
  JOURNAL     Patent: US 5451512-A 68 19-SEP-1995;
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seq_documentation_block:
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  ACCESSION  II4592
  VERSION     II4592.1 GI:997075
  KEYWORDS   Unknown.
  SOURCE      Unknown.
  ORGANISM    Unknown.
  UNCLASSIFIED
  REFERENCE   1 (bases 1 to 270)
  AUTHORS     Apple, R.J., Bugawan, T.L. and Erlich, H.A.
  TITLE       Methods and reagents for HLA class I A locus DNA typing
  JOURNAL     Patent: US 5451512-A 69 19-SEP-1995;
  FEATURES    Location/Qualifiers
  Source      1..270
            /organism="unknown"
BASE COUNT  55 a   84 c   95 g   36 t
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alignment_scores:
  Quality: 44.00      Length: 10
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  Percent Similarity: 100.000  Percent Identity: 90.000

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OM of: US-08-653-294-12 to: N_Geneseq_36.* out_format : pfs
Date: Feb 8, 2000 1:27 PM
About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:
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-O=/cgn1.1/USPTO.spool/US08653294/runat_04022000.160701.15807/app_query.fasta.1
-DB=N_Geneseq_36 -QFMT=fastap -SUFFIX=ring -GAPOP=12.000
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-TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR SCORE=pct
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Search information block:

Query: US-08-653-294-12
Query length: 10
Database: N_Geneseq_36.*
Database sequences: 311585
Database length: 125096042
Search time (sec): 590.520000

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N_Geneseq_36:Q01822	+	44.00	150.12	0.9226	1086	Sequence encoding HLA-Bw52 anti
N_Geneseq_36:Q05693	+	44.00	150.10	0.9255	1089	HLA-B51 gene for production of
N_Geneseq_36:Q05701	+	44.00	150.10	0.9255	1089	HLA-Bw52 gene for production of
N_Geneseq_36:Q12114	+	44.00	150.10	0.9255	1089	HLA-Bw53 exon. HLA-Bw53 gene,
N_Geneseq_36:W70935	+	39.00	131.63	9.89	1026	Sequence encoding the human h
N_Geneseq_36:W70925	+	39.00	120.05	43.68	3874	Sequence of genomic DNA encodi
N_Geneseq_36:W71639	+	39.00	115.47	78.59	6553	HLA B27 consensus sequence. De
N_Geneseq_36:W51732	+	35.00	116.85	65.82	978	DNA encoding a human secreted H
N_Geneseq_36:W33945	+	35.00	110.38	151.02	2056	Human HCMV Inducible gene, SEQ
N_Geneseq_36:Q03811	+	34.00	126.17	19.91	217	Turkey herpes virus (HVT) homol
N_Geneseq_36:W79497	+	34.00	112.63	113.03	1026	Meripilus giganteus galactanase
N_Geneseq_36:W20647	+	34.00	111.68	127.78	1145	Polynucleotide sequence from t
N_Geneseq_36:W35645	+	34.00	105.28	290.30	2386	Cladosporium oxysporum glucose
N_Geneseq_36:W13670	+	33.00	108.49	192.22	1067	Enterococcus faecalis genome c
N_Geneseq_36:W34093	+	33.00	105.41	285.47	1520	Mycobacterium species nucleic
N_Geneseq_36:W30773	+	33.00	102.03	440.33	2240	CD40 associated protein (CAP)-
N_Geneseq_36:W31273	+	33.00	101.58	466.56	2359	LMPI associated protein LAP1 g
N_Geneseq_36:W90123	+	33.00	99.73	591.74	2918	Human CRAFL (TRAF-3) cDNA. Prc
N_Geneseq_36:Q24977	+	33.00	94.79	1.1e+03	5140	DNA encoding soluble mannose
N_Geneseq_36:W30458	+	33.00	68.10	3.3e+04	110000	Rhizobium species plasmid H
N_Geneseq_36:W30459	+	33.00	68.10	3.3e+04	110000	Rhizobium species symbiotic
N_Geneseq_36:W30161	-	32.00	105.24	291.65	1002	Human secreted protein gene 17
N_Geneseq_36:W84507	-	32.00	103.92	345.82	1167	Human secreted protein gene 97
N_Geneseq_36:W07101	+	32.00	102.84	396.88	1320	Staphylococcus aureus mutant H
N_Geneseq_36:W97686	-	32.00	85.41	3.7e+03	9757	Infectious rubella virus RNA.
N_Geneseq_36:W89642	-	32.00	85.41	3.7e+03	9759	Infectious rubella virus cDNA
N_Geneseq_36:W34766	-	32.00	85.41	3.7e+03	9759	Rubella virus RA27/3 genomic s
N_Geneseq_36:W74414	-	32.00	82.23	5.6e+03	14051	Staphylococcus aureus contig
N_Geneseq_36:W21209.07	+	32.00	64.30	5.3e+04	110000	Continuation (8 of 17) of
N_Geneseq_36:W21209.13	+	32.00	64.30	5.3e+04	110000	Continuation (14 of 17) of
N_Geneseq_36:W76405	+	31.00	114.94	84.10	213	Human genome fragment. (Prefe
N_Geneseq_36:W09204	+	31.00	112.04	121.95	297	Virulence factor sequence taker
N_Geneseq_36:W50356	+	31.00	102.89	394.47	849	Sequence encoding fused antibod
N_Geneseq_36:W37348	+	31.00	102.11	435.71	928	Streptococcus pneumoniae coding
N_Geneseq_36:W05843	+	31.00	102.06	438.86	934	Repeat sequence of cps gene loc
N_Geneseq_36:W80055	-	31.00	101.37	479.49	1011	Beta-glucuronidase-contig inser
N_Geneseq_36:W00916	-	31.00	101.37	479.49	1011	Promoter sequence and N-termi
N_Geneseq_36:W01865	+	31.00	100.52	534.40	1114	Fc(epsilon) CH2'-CH4 coding se
N_Geneseq_36:W18352	+	31.00	100.16	559.67	1161	Human V28 seven transmembrane

N_Geneseq_36:Q66170 + 31.00 100.15 560.20 1162 ! Seven transmembrane recepto
N_Geneseq_36:V21328 + 31.00 99.28 626.34 1284 ! Human C epsilon exon. New i
N_Geneseq_36:Q87474 + 31.00 99.12 639.44 1308 ! Human IgE Fc chain (amino a
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seq_documentation_block:

ID Q29167 standard; DNA; 270 BP.
AC Q29167;
DT 09-MAR-1993 (first entry)
DE HLA-Bw 52 exon 2 alpha-1 domain.
KW Human leukocyte antigen; transgenic; germ cells; somatic cells;
expression; ss.
PN J04091731-A.
PD 25-MAR-1992.
PF 03-AUG-1990; 207329.
PR 03-AUG-1990; JP-207329.
PA (OLYU) OLYMPUS OPTICAL CO.
DR WPI; 92-342893/42.
PT Transgenic non-human mammalian HLA-Bw 52 gene - useful for
analysis of expression of gene structure, and prodn. of
mouse model of human disease
PS Disclosure; Fig 1; 8pp; Japanese.
CC The sequence shows the exon 2 alpha-1-domain of the human leukocyte
antigen-Bw 52 gene. The complete gene may be introduced into non-
human mammals, pref. rat or mouse, or their ancestors at the primary
developmental biological step via transplacental into the zygote or
embryo to generate transgenic non-human mammals incorporating the
HLA-Bw 52 gene in both their germ cells and somatic cells. Transgenic
non-human mammals contg. HLA-Bw 52 are useful for the analysis of
expression of the gene, its structure, and prodn. of mouse models of
human disease. See also Q29166-72.
SQ Sequence 270 BP; 59 A; 88 C; 86 G; 37 T;

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Quality: 44.00 Length: 10
Ratio: 4.400 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-12 x Q29167 ..

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seq_name: N_Geneseq_36:Q01834

seq_documentation_block:

ID Q01834 standard; DNA; 1086 BP.
AC Q01834;
DT 19-MAR-1991 (first entry)
DE Sequence encoding HLA-B51 antigen.
KW Probe: HLA class I DNA; immunogen; ss.
OS Homo sapiens.
PN EP35480-A.
PD 14-FEB-1990.
PF 10-AUG-1989.
PR 11-AUG-1988; JP-200758.
PA (OLYU) Olympus Optical Co., Ltd.
DR WPI; 90-046289/07.
PT New DNA for class 1 human leukocyte antigens and derived probes and
transformed cells, useful for DNA typing, as immunogens etc.
PS Claim 1: Page 11; 23pp; English.
CC The HLA class I DNA can be used as a source of probes for use in DNA
typing. Transformed cells, which are useful as immunogens, can be
obtained by introducing these DNAs into eucaryotic cells.
SQ Sequence 1086 BP; 224 A; 334 C; 356 G; 172 T;


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FT /number=5
FT 1013..1042
FT /*tag= f
FT /number=6
FT 1043..1089
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FT 22-AUG-1990.
FT PF 07-FEB-1990; 102424.
FT PR 08-FEB-1989; JP-029313.
FT PA (OLYU ) OLYMPUS OPTICAL KK.
FT PI Takiguchi M.
FT DR WPI; 90-255479/34.
FT PT Allotype specific monoclonal anti- HLA antibodies prodn. - using
FT PT hybridomas derived from transgenic animals carrying HLA gene and
FT PT immunised with HLA antigen of different allele
FT PS Disclosure: Fig 1 A-G; 20pp; English.
FT CC The human HLA-Bw52 gene was introduced into mouse L cells and
FT CC then these cells used to immunise one of the transgenic mice
FT CC (See 005693).
FT CC The spleen lymphocytes were fused with myeloma cells (P3x63-Ag8.653).
FT CC Hybridomas producing antibodies were selected.
FT SQ Sequence 1089 BP; 223 A; 336 C; 359 G; 171 T;

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Ratio: 4.400 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 90.000

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ID Q12114 standard; DNA; 1089 BP.
AC Q12114;
DT 29-AUG-1991 (first entry)
DE HLA-Bw53 exon.
KW Human leukocyte antigen; probe; major histocompatibility complex;
KW MHC; class I; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT cds 1..1089 /*tag= a
FT J03112487-A.
FT PD 14-MAY-1991.
FT PF 22-SEP-1989; 247697.
FT PR 22-SEP-1989; JP-247697.
FT PA (OLYU ) OLYMPUS OPTICAL KK.
FT DR WPI; 91-182991/25.
FT DR P-PSDB; R12463.
FT PT HLA-Bw53 gene, DNA probe and transformant cells - used for
FT immunisation, identifying specificity of antiserum etc.

```

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PS Claim 1; Page 1; 11pp; Japanese.
CC Probes comprising part of the sequence can be used to identify
CC Class I genes. The DNA can be expressed for immunisation of
CC animals and prodn. of monoclonal antibodies specific for the
CC HLA-Bw53 antigen. See also J03112485 and J03112486.
CC Sequence 1089 BP; 222 A; 337 C; 356 G; 174 T;
SQ

alignment_scores:
Quality: 44.00 Length: 10
Ratio: 4.400 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-12 x Q12114 ..
Align seg 1/1 to: Q12114 from: 1 to: 1089

1 ArgGluAspLeuArgIleAlaLeuArgTyr 10
295 CGAGAGAACCTGCGGATCGCGCTCCGCTAC 324

seq_name: N_Geneseq_36:N70935

seq_documentation_block:
ID N70935 standard; DNA; 1026 BP.
AC N70935;
DT 10-APR-1991 (first entry)
DE Sequence encoding the human histocompatibility antigen HLA B27.
DE Rheumatic disorder; genetic screening; diagnosis;
KW ankylosing spondylitis; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT cds 1..1026
FT PN DE3542024-A.
FT PD 04-JUN-1987.
FT PF 28-NOV-1985; 542024.
FT PR 28-NOV-1985; DE-542024.
FT PR 21-DEC-1985; DE-545576.
FT PA (BEHW ) BEHRINGERWERKE AG.
FT PI Riethmuller G, Mec T, Weiss E, Szots H;
FT DR WPI; 87-157893/23.
FT DR P-PSDB; P70590.
FT PT DNA coding for antigen HLA B27 - and diagnostic reagents contg.
FT PS Claim 2; Page 4; 5pp; German.
CC The DNA may be used as a hybridisation probe for detecting the HLA
CC B27 gene, e.g. for assessing susceptibility to rheumatic disorders
CC such as ankylosis spondylitis, or may be used to transform cells
CC for prodn. of HLA B27. The HLA B27 may be used to detect HLA B27
CC antibody in human serum, or to produce mono- or polyclonal HLA B27
CC antibodies for use in immunoassay.
CC Sequence 1026 BP; 213 A; 307 C; 344 G; 162 T;
SQ

alignment_scores:
Quality: 39.00 Length: 10
Ratio: 4.875 Gaps: 0
Percent Similarity: 80.000 Percent Identity: 80.000

alignment_block:
US-08-653-294-12 x N70935 ..
Align seg 1/1 to: N70935 from: 1 to: 1026

1 ArgGluAspLeuArgIleAlaLeuArgTyr 10
223 CGAGAGAACCTGCGGACCTGCTCCGCTAC 252

seq_name: N_Geneseq_36:N70225

seq_documentation_block:
ID N70225 standard; DNA; 3874 BP.

```


This sequence represents a human gene of the invention, that is induced to express by both HCMV and interferon (IFN), designated HCMV-inducible genes (cig or cigs). The invention also relates to genes that are repressed in the presence of HCMV infection, designated HCMV-repressible genes (crg or crgs). The products can be used to obtain agents which can be used for anti-viral therapy, particularly anti-HCMV therapy. They can also be used for the development of drugs that would allow for higher dosage IFN treatments without the concomitant toxicity normally associated with administering high levels of IFN. The products can also be used for detection, diagnosis and drug screening.

FT	sig_peptide	1.	.54
----	-------------	----	-----

```

FT      mat_peptide      /*tag= a
FT      55..1026
FT      /*tag= b
FT      /product= galactanase
FT      /EC_number= 3.2.1.189
PN      WO9732013-A1.
PD      04-SEP-1997.
PF      28-FEB-1997: DK0091.
PR      01-MAR-1996: DK-000234.
PA      (NOVO ) NOVO-NORDISK AS.
PI      Andersen LN, Clausen IG, Kauppinen MS, Kofod LV;
DR      WPI: 97-448685/41.
PT      P-PSDB: W23140.
PS      DPA sequence encoding fungal galactanase - useful in production of
PT      wine or modification of animal feed, e.g. depectinisation, and
PT      reducing viscosity of plant cell wall derived material
PS      Claim 1: Pages 34-36; 49pp; English.
CC      This cDNA sequence encodes a novel galactanase isolated from a
CC      M. giganteus library constructed in E. coli. Positive cDNA inserts were
CC      identified on SC-agar plates using the AZCL xylan assay. cDNA inserts
CC      were amplified directly from yeast colonies. In order to express the
CC      galactanase in Aspergillus, the DNA was digested with appropriate
CC      restriction enzymes, size fractionated on a gel and a fragment
CC      corresponding to the galactanase gene purified. The gene was
CC      subsequently ligated into pHD414 and digested with appropriate
CC      restriction enzymes to give pAZG55. After amplification of the DNA in
CC      E. coli the plasmid was transformed into Aspergillus oryzae and
CC      transformant activity was analysed. The galactanase can be used to
CC      reduce the viscosity of plant cell wall derived material, thus having
CC      implications in wine production, the preparation of fruit or vegetable
CC      juice or for the modification of animal feed allowing a significant
CC      improvement in the in vivo breakdown of plant cell wall material e.g.
CC      depectinisation.
SQ      Sequence 1026 BP; 214 A; 331 C; 285 G; 196 T;

alignment_scores:
  Quality: 34.00      Length: 9
  Ratio: 4.250      Gaps: 0
Percent Similarity: 88.889      Percent Identity: 66.667

alignment_block:
US-08-653-294-12 x T79497/rev ..
Align seg 1/1 to reverse of: T79497 from: 1 to: 1026

      2 GluAspLeuArgIleAlaLeuArgTyr 10
      |||||
564 GAGATCTTCGGTTCCTGCGCTTT 538

seq_name: N_Geneseq_36:X20647

seq_documentation_block:
ID X20647 standard; DNA; 1145 BP.
AC X20647;
DT 05-MAY-1999 (first entry)
DE Polynucleotide sequence from the genome of Treponema pallidum.
KW Treponema pallidum infection; syphilis; Borrelia infection; animal;
KW enzyme production; ds.
OS Treponema pallidum.
PN WO9859034-A2.
PD 30-DEC-1998.
PF 23-JUN-1998: U13041.
PR 24-JUN-1997: US-050667.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Fraser CM;
DR WPI: 99-081273/07.
PT New isolated Treponema pallidum nucleic acids - used to develop
PT products for the detection, diagnosis, characterisation, prevention
PT and therapy of T. pallidum infections, particularly syphilis
PS Claim 1: Page 799-800: 1150pp; English.
CC X20500-21243 represent polynucleotide sequences from the genome of
CC Treponema pallidum. The sequences can be used for detection,

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CC diagnosis, characterisation, prevention and therapy for T. pallidum
CC infections, particularly syphilis. They can also be used for detecting
CC diseases related to Borrelia infections in animals, and for the
CC production of biosynthetic products such as enzymes. 257 T;
SQ      Sequence 1145 BP; 242 A; 267 C; 374 G; 257 T;

alignment_scores:
  Quality: 34.00      Length: 9
  Ratio: 4.250      Gaps: 0
Percent Similarity: 88.889      Percent Identity: 77.778

alignment_block:
US-08-653-294-12 x X20647/rev ..
Align seg 1/1 to reverse of: X20647 from: 1 to: 1145

      2 GluAspLeuArgIleAlaLeuArgTyr 10
      ::|||
839 AAAGATCTTAGGATAGCTCTTGGGTAC 813

seq_name: N_Geneseq_36:V35645

seq_documentation_block:
ID V35645 standard; DNA; 2386 BP.
AC V35645;
DT 08-SEP-1998 (first entry)
DE Cladosporium oxysporum glucose oxidase encoding DNA.
KW Glucose oxidase; Cladosporium oxysporum; enzyme; bread improver;
KW antimicrobial agent; toothpaste; detergent; stain removal;
KW dough additive; hydrogen peroxide generator; ss.
OS Cladosporium oxysporum.
PH Key Location/Qualifiers
FT 5'UTR 1..350
FT /*tag= a
FT CDS 351..2192
FT /*tag= b
FT /product= "glucose oxidase"
FT sig_peptide 351..419
FT /*tag= c
FT /note= "predicted signal sequence"
FT mat_peptide 420..2189
FT /*tag= d
FT 3'UTR 2190..2386
FT /*tag= e
PN WO9820136-A1.
PD 14-MAY-1998.
PF 03-NOV-1997: U20174.
PR 07-NOV-1996; US-746257.
PA (NOVO ) NOVO NORDISK BIOTECH INC.
PI Berka RM, Cherry JR, Halkier T;
DR WPI: 98-286952/25.
DR P-PSDB: W60593.
PT New nucleic acid encoding glucose oxidase active at acidic pH, from
PT Cladosporium - and related vectors and host cells, producing
PT enzyme useful as bread improver, antimicrobial additive for
PT toothpaste, etc. and hydrogen peroxide generator in detergents
PS Claim 4: Fig 2A-B; 83pp; English.
CC This DNA encodes a Cladosporium oxysporum glucose oxidase. Host cells
CC containing a construct comprising the glucose oxidase encoding nucleic
CC acid sequence with regulatory sequences are used to produce recombinant
CC glucose oxidase. Preferred fragments of the nucleic acid are present in
CC pGOX4A and pGOX6A, contained in E. coli deposited as NRRL B-21628 and
CC B-21629. The host cells are particularly Fusarium. The recombinant
CC glucose oxidase is useful as dough additive to improve the gluten
CC quality. It can be used as additive for toothpaste (particularly used
CC with a thiocyanate and lactoperoxidase to generate antimicrobial
CC oxythiocyanate anion), mouthwash, denture cleaners, soaps, hair and body
CC care products. It can be added to cleaning solutions for contact lenses
CC as bleach or as an antibacterial agent. It is also useful as a hydrogen
CC peroxide generator in laundry and dishwashing detergents, particularly
CC for stain removal.

```


SQ Sequence 2386 BP; 583 A; 707 C; 581 G; 515 T;

alignment_scores:
Quality: 34.00 Length: 9
Ratio: 4.250 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 77.778

alignment_block:
US-08-653-294-12 x V35645 ..

Align seg 1/1 to: V35645 from: 1 to: 2386

1 ArgGluAspLeuArgIleAlaLeuArg 9
|||||
605 CGTGAGGACTTACGGACAGCCTTCGA 631

THIS PAGE BLANK (USPTO)

Euthria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 255)
 Hillier L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B.,
 Chisoe, S., Dietrich, N., Dubuque, T., Favello, A., Gish, W.,
 Hawkins, M., Hultman, M., Kucaba, T., Lacy, M., Le, N.,
 Maridis, E., Moore, B., Morris, M., Parsons, J., Prange, C., Rifkin, L.,
 Rohlfing, T., Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J.,
 Trevisan, E., Underwood, K., Wohlmann, P., Waterston, R., Wilson, R.,
 and Marra, M.
 Generation and analysis of 280,000 human expressed sequence tags
 Genome Res. 6 (9), 807-828 (1996)
 9704478
 On May 8, 1995 this sequence version replaced gi:800234.
 Contact: Willson RK
 Washington University School of Medicine
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: est@watson.wustl.edu
 This clone is available royalty-free through LLNL; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 Trace considered overall poor quality
 Seq primer: -28M13 rev2 from Amersham
 High quality sequence stop: 1.
 Location/Qualifiers
 1..255
 /organism="Homo sapiens"
 /db_xref="GDB:4590888"
 /db_xref="taxon:9606"
 /clone_lib="IMAGE:566435"
 /clone_host="Stratagene colon (#937204)"
 /notes="Organ: colon; Vector: pBluescript SK-; Site: 1:
 EcoRI; Site: 2: XhoI; Cloned unidirectionally. Primer:
 Oligo dt: T-84 colonic epithelial cell line. Average
 insert size: 1.0 kb; Uni-ZAP XR Vector; -5' adaptor
 sequence: 5' GAATTCGGCAGG 3' -3' adaptor sequence: 5'
 CTCGAGTTTTTTTTTTT 3'"

BASE COUNT 57 a 70 c 75 g 44 t
 ORIGIN
 alignment_scores
 Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block
 US-08-653-294-10 x AA151891 ..
 Align seg 1/1 to: AA151891 from: 1 to: 255
 1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
 |||||
 77 CGAGAGAACCTGCGATCGCCTCCGCTAC 106

seq_name: gb_est1:AA263158
 seq_documentation_block:
 LOCUS AA263158 283 bp mRNA EST 02-JUL-1998
 DEFINITION PH0534 KGI-a Lambda Zap Express cDNA library Homo sapiens cDNA 5',
 mRNA sequence.
 ACCESSION AA263158
 VERSION AA263158.1 GI:1898964
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Euthria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 283)
 Claudio, J.O., Liew, C.C., Dempsey, A.A., Cukerman, E., Stewart, A.K.,
 Na, E., Atkins, H.I., Iscove, N.N. and Hawley, R.G.
 Identification of sequence-tagged transcripts differentially

expressed within the human hematopoietic hierarchy
 Genomics 50 (1), 44-52 (1998)
 98292493
 On May 5, 1995 this sequence version replaced gi:797810.
 Contact: Hawley RG
 Oncology Research Laboratories
 The Toronto Hospital
 CRCS-424, 67 College St., Toronto, Ontario M5G 2M1, Canada
 Tel: 416 3403834
 Fax: 416 3403453
 Email: r.hawley@utoronto.ca
 Similar to M58636 MHC class I HLA-B* gene. Clone was randomly
 picked from KGIa primary library.
 Seq primer: 5' GAAATTAACCTCACTAAAGG 3'
 High quality sequence stop: 283.
 Location/Qualifiers
 1..283
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="KGI-a Lambda Zap Express cDNA library"
 /cell_type="promyeloblast"
 /cell_line="KGI-a"
 /note="Vector: Lambda Zap Express (Stratagene); Site: 1:
 EcoRI; Site: 2: XhoI; Unidirectional cloning sites:
 EcoRI-XhoI. mRNA was purified from KGI-a cell line, cDNA
 was synthesized using an XhoI-OligodT linker primer. EcoRI
 adaptors were ligated, followed by digestion with XhoI for
 directional cloning into predigested Lambda Zap Express"

BASE COUNT 64 a 91 c 88 g 40 t
 ORIGIN
 alignment_scores
 Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block
 US-08-653-294-10 x AA263158 ..
 Align seg 1/1 to: AA263158 from: 1 to: 283
 1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
 |||||
 120 CGAGAGAACCTGCGATCGCCTCCGCTAC 149

seq_name: gb_est6:D82221
 seq_documentation_block:
 LOCUS D82221 375 bp mRNA EST 09-FEB-1996
 DEFINITION HUMHBC4626 Human pancreatic islet Homo sapiens cDNA similar to
 HLA-B, mRNA sequence.
 ACCESSION D82221
 VERSION D82221.1 GI:1183739
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Euthria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 375)
 Takeda, J.
 Human pancreatic islet ESTs
 Unpublished (1995)
 JOURNAL
 On Apr 14, 1993 this sequence version replaced gi:785255.
 Contact: Jun Takeda
 Institute for Molecular and Cellular Regulation, Gunma University
 3-39-15 Showa-machi, Maebashi Gunma 371, Japan
 Tel: 272-20-8856
 Fax: 272-20-8896
 Email: jtakeda@sb.gunma-u.ac.jp.
 Location/Qualifiers
 1..375
 /organism="Homo sapiens"

JOURNAL MEDLINE
 COMMENT
 On May 5, 1995 this sequence version replaced gi:797810.
 Contact: Hawley RG
 Oncology Research Laboratories
 The Toronto Hospital
 CRCS-424, 67 College St., Toronto, Ontario M5G 2M1, Canada
 Tel: 416 3403834
 Fax: 416 3403453
 Email: r.hawley@utoronto.ca
 Similar to M58636 MHC class I HLA-B* gene. Clone was randomly
 picked from KGIa primary library.
 Seq primer: 5' GAAATTAACCTCACTAAAGG 3'
 High quality sequence stop: 283.
 Location/Qualifiers
 1..283
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone_lib="KGI-a Lambda Zap Express cDNA library"
 /cell_type="promyeloblast"
 /cell_line="KGI-a"
 /note="Vector: Lambda Zap Express (Stratagene); Site: 1:
 EcoRI; Site: 2: XhoI; Unidirectional cloning sites:
 EcoRI-XhoI. mRNA was purified from KGI-a cell line, cDNA
 was synthesized using an XhoI-OligodT linker primer. EcoRI
 adaptors were ligated, followed by digestion with XhoI for
 directional cloning into predigested Lambda Zap Express"

BASE COUNT 64 a 91 c 88 g 40 t
 ORIGIN
 alignment_scores
 Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block
 US-08-653-294-10 x AA263158 ..
 Align seg 1/1 to: AA263158 from: 1 to: 283
 1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
 |||||
 120 CGAGAGAACCTGCGATCGCCTCCGCTAC 149

seq_name: gb_est6:D82221
 seq_documentation_block:
 LOCUS D82221 375 bp mRNA EST 09-FEB-1996
 DEFINITION HUMHBC4626 Human pancreatic islet Homo sapiens cDNA similar to
 HLA-B, mRNA sequence.
 ACCESSION D82221
 VERSION D82221.1 GI:1183739
 KEYWORDS EST.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Euthria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 375)
 Takeda, J.
 Human pancreatic islet ESTs
 Unpublished (1995)
 JOURNAL
 On Apr 14, 1993 this sequence version replaced gi:785255.
 Contact: Jun Takeda
 Institute for Molecular and Cellular Regulation, Gunma University
 3-39-15 Showa-machi, Maebashi Gunma 371, Japan
 Tel: 272-20-8856
 Fax: 272-20-8896
 Email: jtakeda@sb.gunma-u.ac.jp.
 Location/Qualifiers
 1..375
 /organism="Homo sapiens"

/db_xref="taxon:9606"
 /clone_lib="Human pancreatic islet"
 /note="Vector: Lambda ZAPII; Site_1: Eco RI; Site_2: Xho I; mRNA was prepared from normal adult human islets. cDNA was directionally synthesized from the Xho I in the vector to the EcoRI site. cDNA was size fractionated to remove sequences <1000 bp in size."

BASE COUNT 75 a 124 c 118 g 55 t 3 others

alignment_scores:
 Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-10 x D82221 ..

Align seg 1/1 to: D82221 from: 1 to: 375

1 ArgGUAsnLeuArgIleLeuLeuArgTyr 10

|||||

306 CGAGAGACCTGCGATCGCGCTCCGNTAC 335

seq_name: gb_est10:AA147151

seq_documentation_block:
 LOCUS AA147151 581 bp mRNA EST 05-DEC-1996
 DEFINITION Z032d06.r1 Stratagene colon (#937204) Homo sapiens cDNA clone IMAGE:588587 5' similar to gb:M64740 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, A-24(A-9) A*2402 ALPHA (HUMAN);, mRNA sequence.

ACCESSION AA147151
 VERSION AA147151
 KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 581)

AUTHORS Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B., Chissoe, S., Dietrich, N., Dubuque, T., Favello, A., Gish, N., Hawkins, M., Hultman, M., Kucaba, T., Lacy, M., Le, M., Le, N., Mardis, E., Moore, B., Morris, M., Parsons, J., Prange, C., Rifkin, L., Rohlfing, T., Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J., Trevaskis, E., Underwood, K., Wohlmann, P., Waterston, R., Wilson, R., and Marra, M.

TITLE Generation and analysis of 280,000 human expressed sequence tags
 JOURNAL Genome Res. 6 (9), 807-828 (1996)
 MEDLINE 9704478

COMMENT On Sep 12, 1996 this sequence version replaced gi:1393699.

Contact: Wilton RK

Washington University School of Medicine

4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108

Tel: 314 286 1800

Fax: 314 286 1810

Email: est@watson.wustl.edu

This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information.

Seq primer: -28M3 rev2 from Amersham

High quality sequence stop: 272.

Location/Qualifiers

FEATURES

source

1. .581

/organism="Homo sapiens"

/db_xref="GDB:4620889"

/db_xref="taxon:9606"

/clone="IMAGE:588587"

/clone_lib="Stratagene colon (#937204)"

/lab_host="SOLR cells (kanamycin resistant)"

/note="Organ: colon; Vector: pBluescript SK-; Site_1:

EcoRI; Site_2: XhoI; Cloned unidirectionally. Primer:

Oligo dt. 1-84 colonic epithelial cell line. Average

insert size: 1.0 Kb; Uni-ZAP XR Vector; -5' adaptor

sequence: 5' GAATTCGCACGAG 3' -3' adaptor sequence: 5' CTCGAGTTTTTTTTTTT 3'

BASE COUNT 134 a 162 c 185 g 85 t 15 others

ORIGIN

alignment_scores:
 Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-10 x AA147151 ..

Align seg 1/1 to: AA147151 from: 1 to: 581

1 ArgGUAsnLeuArgIleLeuLeuArgTyr 10

|||||

152 CGAGAGAACCTGCGATCGCTCCGCTAC 181

seq_name: gb_est26:AI359260

seq_documentation_block:
 LOCUS AI359260 618 bp mRNA EST 15-FEB-1999
 DEFINITION qy27b07.x1 NCI-CGAP_Brn23 Homo sapiens cDNA clone IMAGE:2013205 3' similar to gb:D32129 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, AW-66(A-10) A*6601 ALPHA (HUMAN);, mRNA sequence.

ACCESSION AI359260

VERSION AI359260.1 GI:4110881

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 618)

AUTHORS NCI/NINDS-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.

TITLE National Cancer Institute / National Institute of Neurological Disorders and Stroke, Brain Tumor Genome Anatomy Project

(CGAP/BTGA), Tumor Gene Index

UNPUBLISHED (1998)

CONTACT: Robert Strausberg, Ph.D.

Tel: (301) 496-1550

Email: Robert.Strausberg@nih.gov

Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D., Ph.D.

cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

cDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 591 Std Error: 0.00

Seq primer: -400P from Gibco

High quality sequence stop: 458.

FEATURES

source

1. 618

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:2013205"

/clone_lib="NCI-CGAP_Brn23"

/tissue_type="glioblastoma (pooled)"

/lab_host="DH10B"

/note="Organ: brain; Vector: pT713D-Pac (Pharmacia) with a

modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st

strand cDNA was primed with a Not I - Oligo(dT) primer [5'

TGTTACCAATCTGAAGTGGAGCGCGCATATCTTTTTTTTTTTTTTTTTTTT

T 3']; double-stranded cDNA was ligated to Eco RI

adaptors (Pharmacia), digested with Not I and cloned into

the Not I and Eco RI sites of the modified pT713 vector.

Library is normalized, and was constructed by Bento

Soares and M.Fatima Bonaldo."

```

BASE COUNT      128 a      171 c      182 g      137 t
ORIGIN

alignment_scores:
  Quality:      44.00      Length:      10
  Ratio:        4.889      Gaps:        0
  Percent Similarity: 90.000      Percent Identity: 90.000

alignment_block:
US-08-653-294-10 x AI359260/rev ..

Align seg 1/1 to reverse of: AI359260 from: 1 to: 618

1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
|||||
319 CGAGAGAACCTGCGGATCGGCTCGCTAC 290

seq_name: gb_est31:AI696864

seq_documentation_block:
LOCUS      AI696864      748 bp      mRNA      EST      03-JUN-1999
DEFINITION wC74h11.x1 NCI-CGAP_Pan1 Homo sapiens cDNA clone IMAGE:2324421 3'
similar to gb:M28205 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
B-51(B-5) B-5101 ALPHA (HUMAN);, mRNA sequence.
ACCESSION      AI696864
VERSION
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS
1 (bases 1 to 748)
TITLE      NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
JOURNAL
COMMENT      Unpublished (1997)
On Mar 16, 1998 this sequence version replaced gi:2961758.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert.Strausberg@nih.gov
Life Technologies catalog #: 11548-013
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco
High quality sequence stop: 424.
FEATURES
source
1..748
Location/Qualifiers
/db_xref="taxon:9606"
/clone="IMAGE:2324421"
/clone_lib="NCI-CGAP_Pan1"
/tissue_type="adenocarcinoma"
/lab_host="DH10B"
/organism="Homo sapiens"
/note="Organ: pancreas; Vector: pCMV-SPOK6; Site 1: SalI;
Site 2: NotI; Cloned unidirectionally. Primer: Oligo dt.
Average insert size 1.72 kb. Life Technologies catalog #:
11548-013"
BASE COUNT      169 a      227 c      237 g      108 t      7 others
ORIGIN

alignment_scores:
  Quality:      44.00      Length:      10
  Ratio:        4.889      Gaps:        0
  Percent Similarity: 90.000      Percent Identity: 90.000

alignment_block:
US-08-653-294-10 x AI696864 ..

Align seg 1/1 to: AI696864 from: 1 to: 748

1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
|||||
137 CGAGAGAACCTGCGGATCGGCTCGCTAC 166

seq_name: gb_est13:AA332511

seq_documentation_block:
LOCUS      AA332511      360 bp      mRNA      EST      21-APR-1997
DEFINITION EST36483 Embryo, 8 week I Homo sapiens cDNA 5' end, mRNA sequence.
ACCESSION      AA332511
VERSION
KEYWORDS      EST.
SOURCE      human.
ORGANISM      Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS
1 (bases 1 to 360)
Adams,M.D., Kerlavage,A.R., Fleischmann,R.D., Fuldner,R.A.,
Bult,C.J., Lee,N.H., Kirkness,E.F., Weinstock,K.G., Gocayne,J.D.,
White,O., Sutton,G., Blake,J.A., Brandon,R.C., Man-Wai,C.,
Clayton,R.A., Cline,T.R., Cotton,M.D., Earle-Hughes,J., Fine,L.D.,
Fitzgerald,L.M., Fitchugh,W.M., Fritchman,J.L., Geoghagen,N.S.,
Glodek,A., Gnehm,C.L., Hanna,M.C., Hedblom,E., Hinkie,P.S., Jr.,
Kelley,J.M., Kelley,J.C., Liu,L.-I., Marmaros,S.M., Merrick,J.M.,
Moreno-Palauques,R.F., McDonald,L.A., Nguyen,D.T., Pelligrino,S.M.,
Phillips,C.A., Ryder,S.E., Scott,J.L., Saudek,D.M., Shirley,R.,
Small,K.V., Spriggs,T.A., Utterback,T.R., Weidman,J.F., Li,Y.,
Bednarek,D.P., Cao,L., Cepeda,M.A., Coleman,T.A., Collins,E.J.,
Dimdek,D., Feng,D.-F., Ferrie,A., Fischer,C., Hastings,G.A.,
He,W.W., Hu,J.S., Greene,J.M., Gruber,J., Hudson,P., Kim,A.K.,
Kozak,D.L., Kunsch,C., Hungjun,J., Li,H., Meissner,P.S., Olsen,H.,
Raymond,L., Wei,Y.F., Wing,J., Xu,C., Yu,G.L., Ruben,S.M.,
Dillon,P.J., Fannon,M.R., Rosen,C.A., Haseltine,W.A., Fields,C.,
Praser,C.M. and Venter,J.C.
Initial assessment of human gene diversity and expression patterns
based upon 83 million nucleotides of cDNA sequence
Nature 377 (6547 Suppl), 3-174 (1995)
12140200
COMMENT      On May 18, 1995 this sequence version replaced gi:811192.
Other ESTs: THC18992
Contact: Kerlavage, AR
Bioinformatics
The Institute for Genomic Research
9712 Medical Center Drive, Rockville, MD 20850 USA
Tel: 3018699056
Fax: 3018699423
Email: arkerlav@tigr.org
For clone availability, additional sequence and expression
information related to this EST, please check the TIGR Human Gene
Index (http://www.tigr.org/tdb/hgi/hgi.html)
Seq primer: M13 Reverse.
FEATURES
source
1..360
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="ATCC (inhost):134173"
/db_xref="taxon:9606"
/clone_lib="Embryo, 8 week I"
/dev_stage="embryo, 8 wks"
/note="Organ: Embryo, 8 weeks; Vector: pBluescript SK-;
Site 1: EcoRI; Site 2: XhoI"
BASE COUNT      87 a      134 c      67 g      67 t      5 others
ORIGIN

alignment_scores:
  Quality:      40.00      Length:      9
  Ratio:        4.444      Gaps:        0
  Percent Similarity: 100.000      Percent Identity: 88.889

alignment_block:
US-08-653-294-10 x AA332511 ..

```

sequence.

TITLE
JOURNAL
COMMENT

WashU-NCI human EST Project
Unpublished (1997)
On Jan 17, 1998 this sequence version replaced gi:1899887.
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: -40ml3 fwd. ET from Amersham.

FEATURES
source
1. .414
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="IMAGE:1539586"
/clone_lib="Johnston frontal cortex"
/sex="male"
/tissue_type="pooled frontal lobe"
/dev_stage="adult"
/lab_host="SOLR (kanamycin resistant)"
/note="Organ: brain; Vector: Bluescript SK-; Site_1:
EcoRI; Stanley Neuropathology Consortium
(www.stanleylab.org) brains S-58, S-65, S-67, S-78.
Random + oligo-dr primed into EcoRI site of ZAP II Vector.
Mass excised. Avg insert length 1.9kb. Custom library
provided by Dr. Nancy Johnston [(410) 614-3918,
nlj@welchlink.welch.jhu.edu].

BASE COUNT 80 a 140 c 136 g 58 t
ORIGIN

alignment_scores:
Quality: 39.00 Length: 10
Ratio: 4.875 Gaps: 0
Percent Similarity: 80.000 Percent Identity: 80.000

alignment_block:
US-08-653-294-10 x A1124815 ..
Align seg 1/1 to: A1124815 from: 1 to: 414

1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
|||||
284 CGAGAGACCTGCGCACCGCGCTCGCTAC 313

seq_name: gb-gss15:AQ614213

seq_documentation_block:
LOCUS AQ614213 498 bp DNA GSS 15-JUN-1999
DEFINITION HS_5123_B1_B06.SP6E RPCI-11 Human Male BAC Library Homo sapiens
genomic clone Plate-699 Col-11 Row-D, genomic survey sequence.

ACCESSION AQ614213
VERSION AQ614213.1 GI:5075489
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 498)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
99380589
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618

Fax: (206) 616-3887
Email: jwallaceu.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 699 row: D column: 11
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 498.

FEATURES
source
1. .498
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate-699 Col-11 Row-D"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Genomic sequence of BAC ends"

BASE COUNT 129 a 121 c 98 g 150 t
ORIGIN

alignment_scores:
Quality: 39.00 Length: 9
Ratio: 4.333 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:
US-08-653-294-10 x A0614213 ..
Align seg 1/1 to: A0614213 from: 1 to: 498

2 GluAsnLeuArgIleLeuLeuArgTyr 10
:::|||||
206 CAGAAATCTCGCTGTGTACTGCGGATAT 232

seq_name: gb_est21:AA989542

seq_documentation_block:
LOCUS AA989542 402 bp mRNA EST 02-JUN-1998
DEFINITION am64d02.sl Barstead spleen HPLRB2 Homo sapiens cDNA clone
(HUMAN):1576803 3' similar to gb:L05093 60S RIBOSOMAL PROTEIN L18A
(HUMAN);, mRNA sequence.

ACCESSION AA989542
VERSION AA989542.1 GI:3174906
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 402)
Hillier,L., Allen,M., Bowles,L., Dubuque,T., Geisel,G., Jost,S.,
Krizman,D., Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M.,
Martin,J., Moore,B., Schellenberg,K., Steptoe,M., Tan,F.,
Theising,B., White,Y., Wylie,T., Waterston,R. and Wilson,R.
WashU-NCI human EST project
Unpublished (1997)
On Jan 19, 1998 this sequence version replaced gi:2153091.
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wustl.edu
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Seq primer: -40ml3 fwd. ET from Amersham
High quality sequence stop: 1.

FEATURES
source
1. .402
Location/Qualifiers

Align seg 1/1 to reverse of: AW092686 from: 1 to: 440

1 ArgGluAsnLeuArgIleLeuLeuArgTyr 10
|||||
142 AGGAGAAATTAGTAATTCTGTACGTTAT 113

Result No.	Score	Query		Length	DB	ID	Description
		Match	%				
1	49	100.0	10	1	W47268	Immunomodulatory p	
2	44	89.8	10	1	W47266	Immunomodulatory p	
3	44	89.8	10	1	W47272	Immunomodulatory p	
4	44	89.8	20	1	R92909	HLA-B*2702 CTL modu	
5	44	89.8	20	1	R92911	HLA-B*2702 CTL modu	
6	44	89.8	20	1	R92907	HLA-B*2702 CTL modu	
7	44	89.8	20	1	R95428	HLA-B*2702 84-75-84	
8	44	89.8	20	1	W33778	Immunomodulating d	
9	44	89.8	20	1	W33779	Immunomodulating d	
10	44	89.8	20	1	W33792	Peptide B2702.84-7	
11	39	79.6	10	1	W47270	Immunomodulatory p	
12	39	79.6	20	1	R92910	HLA-B*2702 CTL modu	
13	39	79.6	20	1	R92908	HLA-B*2702 CTL modu	
14	39	79.6	20	1	R95430	HLA-B*2702 84-75/77	
15	39	79.6	20	1	W33791	Peptide B2702.84-7	
16	39	79.6	20	1	W33793	Peptide B2702.84-7	
17	34	69.4	239	1	W74405	S. aureus gIcB pro	
18	34	69.4	239	1	W74406	S. aureus gIcB pro	
19	34	69.4	383	1	W41592	Rat FRAG1 protein.	
20	34	69.4	485	1	R20796	EHV-4 9C. Nucleic	
21	32	65.3	230	1	R63199	Temp. sensitive au	
22	31	63.3	3079	1	R59926	GAP protein Iraz.	
23	30	61.2	495	1	W89551	Mouse Smad6 protei	
24	30	61.2	496	1	W96815	Smad7 protein used	
25	30	61.2	721	1	W34454	Racillus subtilis	
26	30	61.2	746	1	W34455	Racillus subtilis	
27	30	61.2	851	1	R41333	113 kD ISGF-3alpha	
28	30	61.2	851	1	R72077	Recognition factor	
29	30	61.2	851	1	W03166	Human STAT2. New S	
30	29	59.2	6	1	W47264	Immunomodulatory p	
31	29	59.2	6	1	W33783	Peptide #4 used in	
32	29	59.2	81	1	R62906	TCR alpha chain of	
33	29	59.2	90	1	Y11975	Human 5' EST seque	
34	29	59.2	92	1	R62904	TCR alpha-chain va	

/note= "at least one of the amino acids is the D-isomer"

FT WO9744052-A1.
 FT 27-NOV-1997.
 PD 23-APR-1997; US-651650.
 PR 22-MAY-1996; US-651650.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI; 98-018220/02.
 PT Novel immunomodulatory peptide-type compound - useful for inhibiting
 PT transplant rejection
 PS Claim 10; Page 36; 41pp; English.
 CC The present sequence is an immunomodulatory peptide, which
 CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
 CC in a pharmaceutical composition together with a subtherapeutic dose
 CC of an immunosuppressant, to extend the period of acceptance of a
 CC transplant from a major histocompatibility complex (MHC) unmatched
 CC donor, i.e. to inhibit transplant rejection. It can also be used in
 CC the treatment of autoimmune diseases.
 CC Peptides using the D-form amino acids are more effective
 CC immunomodulators than their diastereomers or enantiomers.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0025;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIIRLNER 10
 DB 1 YRLAIRLNER 10

RESULT 3

WA7272
 ID W47272 standard; peptide; 10 AA.
 AC W47272;
 DT 22-MAY-1998 (first entry)
 DE Immunomodulatory peptide.
 KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
 KW transplant rejection; treatment; autoimmune disease.
 OS Homo sapiens.
 OS Synthetic.
 FH Key Location/Qualifiers
 FT Misc_difference 1..10
 FT /note= "at least one of the amino acids is the D-isomer"

PN WO9744052-A1.
 PD 27-NOV-1997.

PF 23-APR-1997; US-651650.
 PR 22-MAY-1996; US-651650.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI; 98-018220/02.
 PT Novel immunomodulatory peptide-type compound - useful for inhibiting
 PT transplant rejection
 PS Claim 10; Page 36; 41pp; English.
 CC The present sequence is an immunomodulatory peptide, which
 CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
 CC in a pharmaceutical composition together with a subtherapeutic dose
 CC of an immunosuppressant, to extend the period of acceptance of a
 CC transplant from a major histocompatibility complex (MHC) unmatched
 CC donor, i.e. to inhibit transplant rejection. It can also be used in
 CC the treatment of autoimmune diseases.
 CC Peptides using the D-form amino acids are more effective
 CC immunomodulators than their diastereomers or enantiomers.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0025;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLIIRLNER 10
 DB 1 YRLAIRLNER 10

RESULT 4

R92909
 ID R92909 standard; peptide; 20 AA.
 AC R92909;
 DT 16-MAY-1996 (first entry)
 DE HLA-B*2702 CTL modulating peptide (B2702.84-75/75-84(T)).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B*2702.
 OS Synthetic.
 PN WO9526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; US-651650.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI; 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B*75-84 MHC antigen of the recipient
 PT host
 PS Example 15; Page 36; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B*2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIIRLNER 10
 DB 1 YRLAIRLNER 10

RESULT 5

R92911
 ID R92911 standard; peptide; 20 AA.
 AC R92911;
 DT 16-MAY-1996 (first entry)
 DE HLA-B*2702 CTL modulating peptide (B2702.84-75/84-75).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B*2702.
 OS Synthetic.
 PN WO9526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; US-651650.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI; 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B*75-84 MHC antigen of the recipient
 PT host
 PS Example 15; Page 36; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is a dimer of residues 75-84 of the alpha-1 domain of the class I MHC.

CC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054; 1; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 1;
 QY 1 YRLRLRLNER 10
 Db 1 YRLRLRLNER 10
 III IIIIIII
 RESULT 6
 R92907
 ID R92907 standard; peptide; 20 AA.
 AC R92907;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.84-75/75-84).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW Immunosuppressant; graft versus host disorder; transplantation; therapy;
 QS Class I MHC; HLA-B2702.
 OS Synthetic.
 PN W09526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI; 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B7-84 MHC antigen of the recipient
 PT host
 PS Example 15: Page 36; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054; 1; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 1;
 QY 1 YRLRLRLNER 10
 Db 1 YRLRLRLNER 10
 III IIIIIII
 RESULT 7
 R95428
 ID R95428 standard; peptide; 20 AA.
 AC R95428;
 DT 12-NOV-1996 (first entry)
 DE HLA-B2702 84-75-84 palindromic.
 KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
 KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
 KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;

KW cytolysis; antigen presenting cell.
 OS Synthetic.
 PN W09513288-A1.
 PD 18-MAY-1995.
 PF 10-NOV-1994; U12985.
 PR 10-NOV-1993; US-150493.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI; 95-194027/25.
 PT Compns. comprising lymphoid surface membrane proteins - which may
 PT inhibit cytolytic activity and differentiation of CTLs.
 PS Example; Page 12; 29pp; English.
 CC R95413, and R95415-R95431 represent palindromes and fragments of
 CC human-leucocyte-associated antigens. This sequence represents the
 CC HLA-B2702 84-75-84 palindromic. These sequences can be used to isolate
 CC the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
 CC protein associated with T-cell activation in mammalian T-cells, and is
 CC also immunologically cross reactive with the heat shock protein Hsc70.
 CC p74 is found in a limited number of cell types, but is particularly
 CC expressed on B and T cells. p74 can be isolated by lysis of a suitable
 CC cell with an amphoteric detergent, and then passed through an affinity
 CC column containing a covalently bound HLA-B2702 palindromic peptide.
 CC Compositions comprising the extracellular fragment of p74 combined with
 CC HLA-B2702.60-84 (see R95416), induces calcium influx, and inhibits
 CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
 CC compounds can be screened for their effect on the cytolytic activity of
 CC T-cells, by combining them with the extracellular portion of p74 and
 CC determining the amount of binding between the candidate compound and p74.
 CC Modulation of CTL activity can be inhibited in a cellular composition
 CC containing T-cells and antigen presenting cells (APCs), by adding to the
 CC mix the extracellular portion of p74, in an amount sufficient to compete
 CC with p74 for the binding of the p74 ligand.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054; 1; Indels 0; Gaps 0;
 Matches 9; Conservative 0; Mismatches 1;
 QY 1 YRLRLRLNER 10
 Db 1 YRLRLRLNER 10
 III IIIIIII
 RESULT 8
 W33778
 ID W33778 standard; peptide; 20 AA.
 AC W33778;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #1.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744331-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI; 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 16; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is
 CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L), (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a

CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alpha1 domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRLNER 10
 ||| |||||

DB 1 YRLRLRLNER 10

RESULT 9

W33779
 ID W33779 standard; peptide; 20 AA.

AC W33779;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #2.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplanted; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.

PF 22-MAY-1997; U08689.

PR 24-MAY-1996; US-653294.

PA (STRD) UNIV LELAND STANFORD JUNIOR.

PI Beulow R, Clayberger C, Krensky AM;

DR WPI; 98-086530/08

PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases

PS Claim 16; Page 35; 41pp; English.

CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is
 CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alpha1 domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRLNER 10

DB 1 YRLRLRLNER 10
 ||| |||||

RESULT 10

W33792
 ID W33792 standard; peptide; 20 AA.

AC W33792;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.84-75/75-84T tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplanted; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.

PF 22-MAY-1997; U08689.

PR 24-MAY-1996; US-653294.

PA (STRD) UNIV LELAND STANFORD JUNIOR.

PI Beulow R, Clayberger C, Krensky AM;

DR WPI; 98-086530/08.

PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases

PS Example 1; Page 19; 41pp; English.

CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha1 domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0054;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRLNER 10
 ||| |||||

DB 1 YRLRLRLNER 10

RESULT 11

W47270
 ID W47270 standard; peptide; 10 AA.

AC W47270;
 DT 22-MAY-1998 (first entry)
 DE Immunomodulatory peptide.
 KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
 KW transplant rejection; treatment; autoimmune disease.
 OS Homo sapiens.
 OS Synthetic.

PH Key Location/Qualifiers

FT Misc_difference 1.10

FT /note- "at least one of the amino acids is the
 FT D-isomer

PN W09744052-A1.

PD 27-NOV-1997.

PF. 23-APR-1997; U06705.
 PR 22-MAY-1996; US-651650.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI: 98-018220/02.
 PT Novel immunomodulatory peptide-type compound - useful for inhibiting
 PT transplant rejection
 PS Claim 10; Page 36; 41pp; English.
 CC The present sequence is an immunomodulatory peptide, which
 CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
 CC in a pharmaceutical composition together with a subtherapeutic dose
 CC of an immunosuppressant, to extend the period of acceptance of a
 CC transplant from a major histocompatibility complex (MHC) unmatched
 CC donor, i.e. to inhibit transplant rejection. It can also be used in
 CC the treatment of autoimmune diseases.
 CC Peptides using the D-form amino acids are more effective
 CC immunomodulators than their diastereomers or enantiomers.
 SQ Sequence 10 AA;

Query Match 79.6%; Score 39; DB 1; Length 10;
 Best Local Similarity 80.0%; Pred. No. 0.027;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRNER 10
 III IIII
 Db 1 YRLAIRLDR 10

RESULT 12
 R92910
 ID R92910 standard; peptide; 20 AA.
 AC R92910;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.84-75(T)/75-84(T)).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW Class I MHC; HLA-B2702.
 OS Synthetic.
 PN W09526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host
 PS Example 15; Page 36; 80pp; English.
 CC R3061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC Class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.059;
 Matches 8; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLRLRNER 10
 III IIII
 Db 1 YRLAIRLDR 10

RESULT 14
 R95430
 ID R95430 standard; peptide; 20 AA.
 AC R95430;
 DT 12-NOV-1996 (first entry)
 DE HLA-B2702 84-75T/75-84T palindromic.
 KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
 KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
 KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
 KW cytolysis; antigen presenting cell.
 OS Synthetic.
 PN W09513288-A1.
 PD 18-MAY-1995.
 PF 10-NOV-1994; U12985.
 PR 10-NOV-1993; US-150493.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI: 95-194027/25.
 PT Compns. comprising lymphoid surface membrane proteins - which may
 PT inhibit cytolytic activity and differentiation of CTLs.
 PS Example; Page 12; 29pp; English.
 CC R95413, and R95415-R95431 represent palindromes and fragments of
 CC human-leucocyte-associated antigens. These sequences can be used to
 CC isolate the protein p74 from a T-cell lysate. p74 is a T-cell surface
 CC membrane protein associated with T-cell activation in mammalian T-cells,
 CC and is also immunologically cross reactive with the heat shock protein
 CC Hsc70. p74 is found in a limited number of cell types, but is
 CC particularly expressed on B and T cells. p74 can be isolated by lysis of

CC a suitable cell with an amphoteric detergent, and then passed through an
 CC affinity column containing a covalently bound HLA-B2702 palindromic
 CC peptide. Compositions comprising the extracellular fragment of p74
 CC combined with HLA-B2702.60-84 (see R95416), induces calcium influx, and
 CC inhibits cytotoxic T lymphocyte (CTL) differentiation or cytotoxicity.
 CC Candidate compounds can be screened for their effect on the cytolytic
 CC activity of T-cells, by combining them with the extracellular portion of
 CC p74 and determining the amount of binding between the candidate compound
 CC and p74. Modulation of CTL activity can be inhibited in a cellular
 CC composition containing T-cells and antigen presenting cells (APCs), by
 CC adding to the mix the extracellular portion of p74, in an amount
 CC sufficient to compete with p74 for the binding of the p74 ligand.
 CC Sequence 20 AA:

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 88.9%; Pred. No. 0.059;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRNE 9
 ||| |||||
 DB 1 YRLRLRNE 9

RESULT 15

W33791 ID W33791 standard; peptide: 20 AA.
 AC W33791:
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.84-75T/75-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-AL.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DI WPI: 98-086530/08
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 CC Sequence 20 AA:

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.059;
 Matches 8; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLRLRNE 10

DB 1 YRLRLRNE 10
 ||| |||||

Search completed: February 8, 2000, 01:29:38
 Job time: 1750 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:22 ; Search time 117.7 Seconds
(without alignments)
4.008 Million cell updates/sec

Title: US-08-653-294-11
Perfect score: 49
Sequence: 1 YRLIIRLN 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : PIR_62.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	36	73.5	395	2 T01392	leucine-rich repea
2	34	69.4	254	2 JC6168	fibroblast growth
3	34	69.4	485	1 B45343	glycoprotein gp13
4	33	67.3	201	2 S64994	probable membra
5	33	67.3	348	2 I37271	cyclicin II - huma
6	33	67.3	468	1 B46114	glycoprotein gp13
7	33	67.3	506	2 S37583	ring finger protei
8	33	67.3	513	1 TVHURF	ret finger protein
9	33	67.3	560	2 I50372	ORF2 - chicken
10	33	67.3	801	4 TVHURE	transforming prote
11	33	67.3	1711	1 A47392	chromodomain-helic
12	32	65.3	151	2 C71113	probable frxA prot
13	32	65.3	183	2 B70714	hypothetical prote
14	32	65.3	221	2 S56263	hypothetical prote
15	32	65.3	239	2 S57158	hypothetical prote
16	32	65.3	267	2 A32122	dolichyl-phosphate
17	32	65.3	330	2 F64877	peptide transport
18	32	65.3	330	2 S39588	peptide transport
19	32	65.3	468	1 VGBEEH	glycoprotein gp13
20	32	65.3	678	2 B70913	probable penicilli
21	32	65.3	1128	2 A49960	bud emergence prot
22	31	63.3	214	2 T05004	hypothetical prote
23	31	63.3	297	2 T12615	ribosomal protein
24	31	63.3	386	2 T12048	ribosomal protein
25	31	63.3	397	2 T00914	leucine-rich repea
26	31	63.3	400	2 S07733	NADH dehydrogenase
27	31	63.3	425	2 T15959	hypothetical prote
28	31	63.3	825	2 S54465	YTA12 protein prec
29	31	63.3	880	1 B3926	DNA-directed RNA p
30	31	63.3	1131	2 T14517	hypothetical prote

31 31 63.3 2108 2 S28417 CDC39 protein - ye
32 31 63.3 3079 1 RGYI12 probable GTPase-ac
33 30 61.2 144 2 A21047 ribosomal mobile e
34 30 61.2 173 2 S27599 hypothetical prote
35 30 61.2 176 2 F70007 hypothetical prote
36 30 61.2 182 1 TVEFR3 transforming prote
37 30 61.2 186 2 G70418 probable thiamine
38 30 61.2 206 1 E64327 H+-transporting AT
39 30 61.2 232 2 D75062 probable flagella-
40 30 61.2 242 2 B69664 N-acetylglucosamin
41 30 61.2 296 2 A33823 ribosomal protein
42 30 61.2 296 2 B33823 ribosomal protein
43 30 61.2 297 1 RSRTL5 ribosomal protein
44 30 61.2 297 2 S55912 ribosomal protein
45 30 61.2 297 2 JCI308 ribosomal protein

ALIGNMENTS

RESULT 1
T01392
leucine-rich repeat protein T419.11 - Arabidopsis thaliana
C:Species: Arabidopsis thaliana (mouse-ear cross)
C:Date: 12-Feb-1999 #sequence_revision 12-Feb-1999 #text_change 24-Mar-1999
C:Accession: T01392
R:Parnell, L.D.; Gnoj, L.; de la Bastide, M.; Hameed, A.; Habermann, K.; Schutz, K.;
submitted to the EMBL Data Library, May 1998
A:Description: Genomic sequence of BAC T419 from Arabidopsis thaliana, chromosome IV,
A:Reference number: Z14314
A:Accession: T01392
A:Status: translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 1-395 <PAR>
A:Cross-references: EMBL:AF069442; NID:g3242970; PID:g3924504
A:Experimental source: cultivar Columbia
C:Genetics:
A:Map position: 4
A:Note: T419.11

Query Match 73.5%; Score 36; DB 2; Length 395;
Best Local Similarity 66.7%; Pred. No. 7.3;
Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLIIRLN 9
DB 231 YRVLLRLNQ 239
||:|:|:|:
|:|:|:|:

RESULT 2
JC6168
fibroblast growth factor receptor activating protein 1 - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 11-Apr-1997 #sequence_revision 09-May-1997 #text_change 10-Sep-1997
C:Accession: JC6168
R:Lorenz, M.V.; Horii, Y.; Yamana, K.; Sakaguchi, K.; Miki, T.
Proc. Natl. Acad. Sci. U.S.A. 93, 8956-8961, 1996
A:Title: FRAG1, a gene that potentially activates fibroblast growth factor receptor by C
A:Reference number: JC6168; MUID:96392347
A:Contents: osteosarcoma cell
A:Accession: JC6168
A:Molecule type: mRNA
A:Residues: 1-254 <LOR>
A:Cross-references: GB:U57715; NID:g1518608; PID:g1518609
A:Note: the authors translated the codon GTG for residue 56 as Cys
C:Comment: This protein plays an important role in cellular functions, and in fibrobl
C:Genetics:
A:Gene: frog1
C:Keywords: growth factor receptor; osteosarcoma

Query Match 69.4%; Score 34; DB 2; Length 254;

Best Local Similarity 87.5%; Pred. No. 12;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 YRLRLN 8
|||||
Db 109 YRLRLN 116

RESULT 3

B45343
glycoprotein gp13 precursor - equine herpesvirus 4
N:Alternate names: glycoprotein 9C
C:Species: equine herpesvirus 4
C:Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 16-Jul-1999
A:Accession: B45343
R:Nicolson, L.; Onions, D.E.
Virology 179, 378-387, 1990
A:Title: The nucleotide sequence of the equine herpesvirus 4 gC gene homologue.
A:Reference number: A45343; MUID:91021040
A:Accession: B45343
A:Molecule type: DNA
A:Residues: 1-485 <NIC>
A:Cross-references: GB:M58031; NID:g330894; PIDN:AAA46083.1; PID:g330896
C:Genetics:
A:Gene: 16
C:Superfamily: herpesvirus glycoprotein F
C:Keywords: glycoprotein; transmembrane protein
F:1-30/Domain: signal sequence #status predicted <SIG>
F:31-485/Product: glycoprotein gp13 #status predicted <GGP>
F:60,61,66,67,72,108,116,147,220,225,286/Binding site: carbohydrate (Asn) (covalent) #st

Query Match 69.4%; Score 34; DB 1; Length 485;
Best Local Similarity 70.0%; Pred. No. 23;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Oy 1 YRLRLN 10
|||||
Db 120 YRLRLN 129

RESULT 4

S64994
probable membrane protein YLR145w - yeast (Saccharomyces cerevisiae)
N:Alternate names: hypothetical protein I3301
C:Species: Saccharomyces cerevisiae
C:Date: 01-Aug-1995 #sequence_revision 24-May-1996 #text_change 26-Aug-1999
A:Accession: S64994
R:Rieger, M.; Mueller-Auer, S.; Brueckner, M.
submitted to the Protein Sequence Database, May 1996
A:Reference number: S64997
A:Accession: S64994
A:Molecule type: DNA
A:Residues: 1-201 <RIE>
A:Cross-references: EMBL:Z73317; NID:g1360569; PID:e245582; PID:g1360570; MIPS:YLR145w
A:Experimental source: strain S288C
C:Genetics:
A:Map position: 12R
C:Superfamily: Saccharomyces probable membrane protein YLR145w
C:Keywords: transmembrane protein
F:92-108/Domain: transmembrane #status predicted <TM>

Query Match 67.3%; Score 33; DB 2; Length 201;
Best Local Similarity 60.0%; Pred. No. 15;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Oy 1 YRLRLN 10
|||||
Db 15 YRLRLN 24

RESULT 5

...

I37271
cylicin II - human
C:Species: Homo sapiens (man)
C:Date: 12-Aug-1996 #sequence_revision 12-Aug-1996 #text_change 25-Oct-1996
A:Accession: I37271; S52774
R:Hess, H.; Held, H.; Zimbelmann, R.; Franke, W.W.
Exp. Cell Res. 218, 174-182, 1995
A:Title: The protein complexity of the cytoskeleton of bovine and human sperm heads:
A:Reference number: I37271; MUID:95255491
A:Accession: I37271
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-348 <HES>
A:Cross-references: EMBL:Z46788; NID:g758586; PID:g758587

Query Match 67.3%; Score 33; DB 2; Length 348;
Best Local Similarity 60.0%; Pred. No. 27;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Oy 1 YRLRLN 10
|||||
Db 78 YRLRLN 87

RESULT 6

B46114
glycoprotein gp13 precursor - equine herpesvirus 1 (strain Kentucky A)
N:Alternate names: glycoprotein C
C:Species: equine herpesvirus 1
A:Note: host Equus caballus (domestic horse)
C:Date: 17-Feb-1994 #sequence_revision 17-Feb-1994 #text_change 16-Jul-1999
C:Accession: B46114
R:Matsumura, T.; Smith, R.H.; O'Callaghan, D.J.
Virology 193, 910-923, 1993
A:Title: DNA sequence and transcriptional analyses of the region of the equine herpes
A:Reference number: A46114; MUID:93212524
A:Accession: B46114
A:Molecule type: DNA
A:Residues: 1-468 <MAT>
A:Cross-references: GB:S57839; NID:g298846; PIDN:AAB25944.1; PID:g298848
C:Superfamily: herpesvirus glycoprotein F
C:Keywords: glycoprotein; transmembrane protein
F:1-30/Domain: signal sequence #status predicted <SIG>
F:31-468/Product: glycoprotein gp13 #status predicted <GPT>
F:432-451/Domain: transmembrane #status predicted <TM>
F:46,57,62,92,100,131,203,208,269/Binding site: carbohydrate (Asn) (covalent) #status

Query Match 67.3%; Score 33; DB 1; Length 468;
Best Local Similarity 70.0%; Pred. No. 36;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Oy 1 YRLRLN 10
|||||
Db 104 YRLRLN 113

RESULT 7

S37583
RING finger protein rfp - mouse (fragment)
C:Species: Mus musculus (house mouse)
C:Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 16-Jul-1999
A:Accession: S37583
R:Takahashi, M.
submitted to the EMBL Data Library, October 1993
A:Reference number: S37583
A:Accession: S37583
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-506 <TAK>
A:Cross-references: EMBL:X75343; NID:g406747; PIDN:CAA53092.1; PID:g406748
C:Superfamily: rfp transforming protein; RING finger homology

C:Keywords: zinc

F:5-55/Domain: RING finger homology <RNG>

Query Match 67.3%; Score 33; DB 2; Length 506;
Best Local Similarity 77.8%; Pred. No. 39;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLIIRLNE 9

||||| ||

Db 191 YRLRLARLEE 199

RESULT 8

TVHURE

N:Alternates: ret finger protein - human

N:Alternate names: transforming protein rfp

C:Species: Homo sapiens (man)

C:Date: 30-Sep-1989 #sequence_revision 30-Sep-1989 #text_change 18-Jun-1999

C:Accession: A28101

R:Takahashi, M.; Inaguma, Y.; Hiai, H.; Hirose, F.

Mol. Cell. Biol. 8, 1853-1856, 1988

A:Title: Developmentally regulated expression of a human "finger"-containing gene encode

A:Reference number: A28101; MUID:88246464

A:Accession: A28101

A:Molecule type: mRNA

A:Residues: 1-513 <TAK>

A:Cross-references: DDBJ:J03407; NID:g337371; PIDN:AAA36564.1; PID:g337372

C:Genetics:

A:Gene: GDB:RFP

A:Cross-references: GDB:511359; GDB:1391662

A:Map position: 6p22-6p21.3

C:Superfamily: rfp transforming protein; RING finger homology

C:Keywords: DNA binding; transforming protein; zinc

F:1-315/Product: transforming protein rfp (fragment) #status predicted <RET>

F:12-62/Domain: RING finger homology <RNG>

F:16-127/Domain: metal and nucleic acid binding #status predicted <TMN>

Query Match

Best Local Similarity 67.3%; Score 33; DB 1; Length 513;

Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLIIRLNE 9

||||| ||

Db 198 YRLRLARLEE 206

RESULT 9

ORF2 - chicken

C:Species: Gallus gallus (chicken)

C:Date: 13-Sep-1996 #sequence_revision 13-Sep-1996 #text_change 22-Jun-1999

C:Accession: I50372

R:Funahashi, J.; Sekido, R.; Murai, K.; Kamachi, Y.; Kondoh, H.

Development 119, 433-446, 1993

A:Title: Delta-crystallin enhancer binding protein delta EF1 is a zinc finger-homeodoma

A:Reference number: I50222; MUID:94116444

A:Accession: I50372

A:Status: preliminary; translated from GB/EMBL/DDBJ

A:Molecule type: mRNA

A:Residues: 1-560 <FUN>

A:Cross-references: GB:D14316; NID:g391639; PIDN:BA03262.1; PID:g391640

C:Superfamily: CHD-1 protein; chromobox homology

Query Match

Best Local Similarity 67.3%; Score 33; DB 2; Length 560;

Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLLIIRLNER 10

||||| ||

Db 43 KLLIIRLNER 51

RESULT 10

TVHURE

transforming protein RFP/protein-tyrosine kinase RET mutant fusion protein - human

N:Alternate names: ret oncogene protein

N:Contains: protein-tyrosine kinase (EC 2.7.1.112) ret

C:Species: Homo sapiens (man)

C:Date: 31-Mar-1989 #sequence_revision 10-Sep-1997 #text_change 13-Aug-1999

C:Accession: A27203

R:Takahashi, M.; Cooper, G.M.

Mol. Cell. Biol. 7, 1378-1385, 1987

A:Title: ret transforming gene encodes a fusion protein homologous to tyrosine kinase

A:Reference number: A27203; MUID:87257826

A:Accession: A27203

A:Molecule type: mRNA

A:Residues: 1-801 <TAK>

A:Cross-references: GB:M16029; NID:g340025

A:Note: Codons preceding the probable start codon were translated

C:Comment: The ret oncogene is the chimeric product of a translocation mutation betwe

C:Genetics:

A:Gene: RFP/RET

C:Keywords: ATP: fusion protein; oncogene; phosphotransferase; transforming protein;

F:1-315/Region: transforming protein rfp

F:316-792/Region: protein-tyrosine kinase ret

F:459-467/Region: protein kinase ATP-binding motif

F:487/Active site: Lys #status predicted

Query Match

Best Local Similarity 67.3%; Score 33; DB 4; Length 801;

Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLIIRLNE 9

||||| ||

Db 198 YRLRLARLEE 206

RESULT 11

A47392

chromodomain-helicase-DNA-binding protein, CHD-1 - mouse

N:Alternate names: KVPB protein

C:Species: Mus musculus (house mouse)

C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999

C:Accession: A47392; S21568

R:Delmas, V.; Stokes, D.G.; Perry, R.P.

Proc. Natl. Acad. Sci. U.S.A. 90, 2414-2418, 1993

A:Title: A mammalian DNA-binding protein that contains a chromodomain and an SNF2/SWI

A:Reference number: A47392; MUID:93211972

A:Accession: A47392

A:Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-1711

A:Experimental source: S194 plasmacytoma cells

A:Note: Sequence inconsistent with the nucleotide translation

R:Delmas, V.; Perry, R.P.

submitted to the EMBL Data Library, May 1992

A:Description: KVPB, a mammalian protein that contains the SNF2/SWI2 helicase domain

A:Reference number: S21568

A:Accession: S21568

A:Molecule type: mRNA

A:Residues: 772-1711 <DE2>

A:Cross-references: EMBL:X66028

C:Superfamily: CHD-1 protein; chromobox homology

C:Keywords: DNA binding

F:293-336/Domain: chromobox homology <CB1>

F:387-427/Domain: chromobox homology <CB2>

Query Match

Best Local Similarity 67.3%; Score 33; DB 1; Length 1711;

Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

C>Date: 02-Sep-1995 #sequence_revision 19-Oct-1995 #text_change 05-Dec-1997
 C:Accession: S56263
 R:Murakami, Y.; Naitou, M.; Hagiwara, H.; Shibata, T.; Ozawa, M.; Sasanuma, S.I.; Sas
 submitted to the EMBL Data Library, May 1995
 A:Description: Analysis of the nucleotide sequence of chromosome VI from *Saccaromyces*
 A:Reference number: S56186
 A:Accession: S56263
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-221 <MUR>
 A:Cross-references: EMBL:D50617; NID:g836685; PID:d1009888; PID:g836763; MIPS:YFR008w
 C:Genetics:
 A:Map position: 6R

Query Match 65.3%; Score 32; DB 2; Length 221;
 Best Local Similarity 75.0%; Pred. No. 27;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLRLRN 8
 |||||
 DB 211 YRLRLRN 218

RESULT 15

S57158
 hypothetical protein YJRL35c - yeast (*Saccharomyces cerevisiae*)
 N:Alternate names: hypothetical protein J2122
 C:Species: *Saccharomyces cerevisiae*
 C>Date: 08-Jul-1995 #sequence_revision 08-Sep-1995 #text_change 06-Feb-1998
 C:Accession: S57158
 R:Rose, M.; Koetter, P.; Entian, K.D.
 submitted to the Protein Sequence Database, September 1995
 A:Reference number: S56848
 A:Accession: S57158
 A:Molecule type: DNA
 A:Residues: 1-239 <ROS>
 A:Cross-references: EMBL:Z49635; NID:g1015871; PID:g1015872; MIPS:YJRL35c
 C:Genetics:
 A:Gene: SGD:MCM22
 A:Cross-references: SGD:S0003896; MIPS:YJRL35c
 A:Map position: 10R

Query Match 65.3%; Score 32; DB 2; Length 239;
 Best Local Similarity 75.0%; Pred. No. 29;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRN 8
 |||||
 DB 205 YRLRLRN 212

Search completed: February 7, 2000, 11:54:23
 Job time: 24333 sec

QY 2 RLLRLNR 10
 |||||
 DB 793 KLLRLNR 801

RESULT 12
 C71113
 probable frxA protein - *Pyrococcus horikoshii*
 C:Species: *Pyrococcus horikoshii*
 C>Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 14-Aug-1998
 C:Accession: C71113
 R:Kawarabayashi, Y.; Sawada, M.; Horikawa, H.; Haikawa, Y.; Hino, Y.; Yamamoto, S.; Seki
 M.; Ohfuku, Y.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; Kushida, N.; Oguchi
 DNA Res. 5, 55-76, 1998
 A>Title: Complete sequence and gene organization of the genome of a hyper-thermophilic
 A:Reference number: A71000; MUID:98344137
 A:Accession: C71113
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-151 <KAW>
 A:Cross-references: GB:AP000003; NID:g3236130; PID:d1030708; PID:g3257082
 A:Experimental source: strain O73
 A>Note: This accession replaces an interim accession for a sequence replaced by GenBank
 C:Genetics:
 A:Gene: PH0674

Query Match 65.3%; Score 32; DB 2; Length 151;
 Best Local Similarity 77.8%; Pred. No. 18;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLLRLNR 10
 |||||
 DB 96 RLLRLNR 104

RESULT 13

B70744
 hypothetical protein Rv0487 - *Mycobacterium tuberculosis* (strain H37RV)
 C:Species: *Mycobacterium tuberculosis*
 C>Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 17-Jul-1998
 C:Accession: B70744
 R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.
 Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 Nature 393, 537-544, 1998
 A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
 A>Title: Deciphering the biology of *Mycobacterium tuberculosis* from the complete genome
 A:Reference number: A70500; MUID:98295987
 A:Accession: B70744
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-183 <COL>
 A:Cross-references: GB:Z77162; GB:AL123456; NID:g3261606; PID:e255029; PID:g1449285
 A:Experimental source: strain H37RV
 C:Genetics:
 A:Gene: Rv0487

Query Match 65.3%; Score 32; DB 2; Length 183;
 Best Local Similarity 60.0%; Pred. No. 22;
 Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLRLNR 10
 |||||
 DB 77 YRLRLNR 86

RESULT 14

S56263
 hypothetical protein YFR008w - yeast (*Saccharomyces cerevisiae*)
 C:Species: *Saccharomyces cerevisiae*

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2000, 00:59:51 ; Search time 63.71 Seconds
(without alignments)
4.688 Million cell updates/sec

Title: US-08-653-294-11
Perfect score: 49
Sequence: 1 YRLIHLNER 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 82229 seqs, 29864866 residues

Total number of hits satisfying chosen parameters: 82229

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : SwissProt_38:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	34	69.4	485	1 VGLC_HSV4	P22596 equine herp
2	33	67.3	299	1 RL5_BOMMO	O76190 bombyx mori
3	33	67.3	348	1 CYL2_HUMAN	Q14093 homo sapien
4	33	67.3	513	1 RFP_HUMAN	P14373 homo sapien
5	33	67.3	522	1 RFP_MOUSE	Q62158 mus musculu
6	33	67.3	1709	1 CHD1_HUMAN	O14646 homo sapien
7	33	67.3	1711	1 CHD1_MOUSE	P40201 mus musculu
8	32	65.3	183	1 Y487_MYCTU	Q11153 mycobacteri
9	32	65.3	221	1 YFH8_YEAST	P43592 saccharomyc
10	32	65.3	239	1 Y9D_YEAST	P47167 saccharomyc
11	32	65.3	267	1 DPM1_YEAST	P14020 saccharomyc
12	32	65.3	327	1 RL5_ANOGA	O44248 anopheles g
13	32	65.3	330	1 SAPD_ECOLI	P36635 escherichia
14	32	65.3	330	1 SAPD_SALTY	P36636 salmonella
15	32	65.3	468	1 VGLC_HSV4	P12889 equine herp
16	32	65.3	1128	1 BEM3_YEAST	P32873 saccharomyc
17	31	63.3	294	1 RL5A_SCHPO	P52822 schizosacch
18	31	63.3	294	1 RL5B_SCHPO	O74306 schizosacch
19	31	63.3	297	1 RL5_HELAN	O65353 helianthus
20	31	63.3	386	1 RL4_URECA	P49165 urechis cau
21	31	63.3	400	1 NUCM_PASTE	P15689 paramecium
22	31	63.3	825	1 RCAL_YEAST	P40341 saccharomyc
23	31	63.3	880	1 RPA1_SULAC	P11512 sulfolobus
24	31	63.3	2108	1 NOT1_YEAST	P25655 saccharomyc
25	31	63.3	3079	1 IRA2_YEAST	P19158 saccharomyc
26	30	61.2	124	1 RL5_PIG	O95276 sus scrofa
27	30	61.2	206	1 ATPE_METJA	O57673 methanococc
28	30	61.2	242	1 NAGB_BACSU	O35000 bacillus su
29	30	61.2	293	1 RL5_CAEEL	P49405 caenorhabd
30	30	61.2	295	1 RL5A_XENLA	P15125 xenopus lae
31	30	61.2	295	1 RL5B_XENLA	P15125 xenopus lae
32	30	61.2	296	1 RL5_CHICK	P22451 gallus gall
33	30	61.2	296	1 RL5_HUMAN	P46777 homo sapien
34	30	61.2	296	1 RL5_RAT	P09895 rattus norv

ALIGNMENTS

```

RESULT_1
VGLC_HSV4
ID VGLC_HSV4 STANDARD; PRT; 485 AA.
AC P22596;
DT 01-AUG-1991 (Rel. 19, Created)
DT 01-AUG-1991 (Rel. 19, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE GLYCOPROTEIN C PRECURSOR (GLYCOPROTEIN 13).
GN GC OR GP13.
OS Equine herpesvirus type 4 (strain 1942) (EHV-4) (Equine herpesvirus
type 1 subtype 2).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 91021040.
RA NICOLSON L., ONIONS D.E.;
RT "The nucleotide sequence of the equine herpesvirus 4 gc gene
homologue."
RL Virology 179:378-387(1990).
CC -1- SIMILARITY: TO OTHER HERPESVIRUSES GLYCOPROTEIN C.
CC -1- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN GENE SUPERFAMILY.
CC -----
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CC -----
DR EMBL; M58031; AAA46083.1; --
DR EMBL; A21044; CAA01528.1; --
DR PIR; B45343; B45343.
KW Glycoprotein; Transmembrane; Signal.
FT SIGNAL 1 32
FT CHAIN 33 485 GLYCOPROTEIN C.
FT DOMAIN 33 444 EXTRACELLULAR.
FT TRANSEM 445 468
FT CARBOHYD 60 60 POTENTIAL.
FT CARBOHYD 61 61 POTENTIAL.
FT CARBOHYD 66 66 POTENTIAL.
FT CARBOHYD 67 67 POTENTIAL.
FT CARBOHYD 72 72 POTENTIAL.
FT CARBOHYD 108 108 POTENTIAL.
FT CARBOHYD 116 116 POTENTIAL.
FT CARBOHYD 147 147 POTENTIAL.
FT CARBOHYD 220 220 POTENTIAL.
FT CARBOHYD 225 225 POTENTIAL.
FT CARBOHYD 286 286 POTENTIAL.
SQ: SEQUENCE 485 AA; 52509 MW; 63F72464 CRC32;

```

Query Match 69.4%; Score 34; DB 1; Length 485;
Best Local Similarity 70.0%; Pred. No. 11;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

```

QY 1 YRLRLRNER 10
   ||| | |||
Db 120 YRLRLRNER 129

RESULT 2
RL5_BOMMO
ID RL5_BOMMO STANDARD; PRT; 299 AA.
AC 076190;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE 60S RIBOSOMAL PROTEIN L5.
GN RPL5.
OS Bombyx mori (Silk moth).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrysia;
OC Bombycoidea; Bombycidae; Bombyx.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=SILK GLAND;
RA YANG C.S., SEHNAL F.;
RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: THIS PROTEIN BINDS 5S RNA (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE L18P FAMILY OF RIBOSOMAL PROTEINS.
CC -----
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CC -----
CC EMBL: AF008229; AAC24960.1;
CC DR PFAM: PF00861; Ribosomal_L18p; 1.
CC KW Ribosomal protein; rRNA-binding.
CC SQ SEQUENCE 299 AA; 34378 MW; 7262D2FC CRC32;

Query Match 67.3%; Score 33; DB 1; Length 299;
Best Local Similarity 50.0%; Pred. No. 10;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRNER 10
   ||| | |||
Db 49 YRLRLRNSK 58

RESULT 3
CYL2_HUMAN
ID CYL2_HUMAN STANDARD; PRT; 348 AA.
AC Q14093;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE CYLICIN II (MULTIPLE-BAND POLYPEPTIDE II).
GN CYL2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=TESTIS;
RA HESS H., HEID H., ZIMMELMANN R., FRANK W.W.;
RX MEDLINE: 95255491.
RT "The protein complexity of the cytoskeleton of bovine and human sperm
RT heads: the identification and characterization of cylicin II."
RL Exp. Cell Res. 218:174-182(1995).
CC -!- FUNCTION: POSSIBLE ARCHITECTURAL ROLE DURING SPERMATOGENESIS. MAY
CC BE INVOLVED IN SPERMATID DIFFERENTIATION.
CC -!- SUBCELLULAR LOCATION: CALYX; SPERM HEAD CYTOSKELETAL STRUCTURE.
CC -!- TISSUE SPECIFICITY: TESTIS.

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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: 246788; CAA86752.1;
CC KW Cytoskeleton; Structural protein; Repeat; Sperm; Spermatogenesis.
CC FT DOMAIN 25 347 31 X 3 AA REPEATS OF K-K-X.
CC FT REPEAT 157 240 3 X APPROXIMATE TANDEM REPEATS.
CC FT REPEAT 157 184 1.
CC FT REPEAT 185 212 2.
CC FT REPEAT 213 240 3.
CC SQ SEQUENCE 348 AA; 39079 MW; FD27FBF CRC32;

Query Match 67.3%; Score 33; DB 1; Length 348;
Best Local Similarity 60.0%; Pred. No. 12;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRNER 10
   || | |||
Db 78 YRLRLRNER 87

RESULT 4
RFP_HUMAN
ID RFP_HUMAN STANDARD; PRT; 513 AA.
AC P14373;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE ZINC-FINGER PROTEIN RFP (RET FINGER PROTEIN).
GN RFP.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 88246464.
RA TAKAHASHI M., INAGUMA Y., HIAI H., HIROSE F.;
RT "Developmentally regulated expression of a human 'finger'-containing
RT gene encoded by the 5' half of the ret transforming gene."
RL Mol. Cell Biol. 8:1853-1856(1988).
CC -!- FUNCTION: MAY FUNCTION IN MALE GERM CELL DEVELOPMENT.
CC -!- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC -!- DISEASE: RECOMBINATION OF THE N-TERMINAL OF RFP WITH A PROTEIN
CC TYROSINE KINASE PRODUCES THE RET TRANSFORMING PROTEIN.
CC -!- SIMILARITY: CONTAINS A C3HC4-CLASS ZINC FINGER.
CC -----
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CC -----
CC EMBL: J03407; AAA36564.1;
CC DR PIR: A28101; TVHURF.
CC DR MIM: 602165;
CC DR PROSITE: PS00518; ZINC_FINGER_C3HC4; 1.
CC DR PFAM: PF00097; Zf-C3HC4; 1.
CC DR PFAM: PF00622; SPRY; 1.
CC DR PFAM: PF00643; Zf-B_Box; 1.
CC KW Proto-oncogene; Zinc-finger; Metal-binding; Chromosomal translocation;
CC Nuclear protein; DNA-binding.
CC SITE 315 316
CC FT SITE 315 316
CC FT BREAKPOINT FOR TRANSLOCATION TO FORM THE
CC RFP-RET ONCOGENE.
CC FT C3HC4-TYPE.

```

FT DOMAIN 96 127 B BOX.
SO SEQUENCE 513 AA; 58489 MW; 022BC859 CRC32;

Query Match 67.3%; Score 33; DB 1; Length 513;
Best Local Similarity 77.8%; Pred. No. 18;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLRLRNE 9
||| |||
Db 198 YRLRLRLEE 206

RESULT 5

REF_MOUSE
ID RFP_MOUSE STANDARD; PRT; 522 AA.
AC O62158;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE ZINC-FINGER PROTEIN RFP (RET FINGER PROTEIN).
GN RFP.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6;
RX MEDLINE: 97176437.
RA CAO T., SHANNON M., HANDEL M.A., ETKIN L.D.;
RT "Mouse ret finger protein (rpf) proto-oncogene is expressed at
RT specific stages of mouse spermatogenesis.";
RL Dev. Genet. 19:309-320(1996).
CC -!- FUNCTION: MAY FUNCTION IN MALE GERM CELL DEVELOPMENT.
CC -!- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC -!- SIMILARITY: CONTAINS A C3HC4-CLASS ZINC FINGER.

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CC -----
CC EMBL: L46855; AAA85354.1; -
CC MGD: MGI:97904; RFP.
CC PROSITE: PS00518; ZINC_FINGER_C3HC4; 1.
CC PFAM: PF00097; zf-C3HC4; 1.
CC PFAM: PF00622; SPRY; 1.
CC PFAM: PF00643; zf-B_box; 1.
CC Zinc-finger; Metal-binding; Nuclear protein; DNA-binding.
FT ZN_FING 25 65 C3HC4-TYPE.
FT DOMAIN 105 136 B BOX.
SQ SEQUENCE 522 AA; 59550 MW; 18E6E716 CRC32;

Query Match 67.3%; Score 33; DB 1; Length 522;
Best Local Similarity 77.8%; Pred. No. 19;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLRLRNE 9
||| |||
Db 207 YRLRLRLEE 215

RESULT 6

CHDI_HUMAN
ID CHDI_HUMAN STANDARD; PRT; 1709 AA.
AC O14646;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)

DE CHROMODOMAIN-HELICASE-DNA-BINDING PROTEIN 1 (CHD-1).
GN CHDI.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Homiidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 97470991.
RA WOODAGE T., BASRAI M.A., BAXEVANIS A.D., HIETER P., COLLINS F.S.;
RT "Characterization of the CHD family of proteins.";
RL Proc. Natl. Acad. Sci. U.S.A. 94:11472-11477(1997).
CC -!- FUNCTION: SEQUENCE-SELECTIVE DNA-BINDING PROTEIN. COULD PLAY AN
CC IMPORTANT ROLE IN GENE REGULATION.
CC -!- SUBCELLULAR LOCATION: NUCLEAR.
CC -!- SIMILARITY: BELONGS TO THE SNF2/RAD54 HELICASE FAMILY.
CC -!- SIMILARITY: CONTAINS 2 'CHROMO' DOMAINS.
CC -----
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CC -----
CC EMBL: AF006513; AAB87381.1; -
CC HSP: P23197; LAP0.
CC MIM: 602118; -
CC DR PROSITE: PS00598; CHROMO_1; 2.
CC DR PROSITE: PS00013; CHROMO_2; 2.
CC DR PFAM: PF00176; SNF2_N; 1.
CC DR PFAM: PF00271; helicase_C; 1.
CC DR PFAM: PF00385; Chromo; 2.
CC KW DNA-binding; ATP-binding; Helicase; Nuclear protein; Repeat.
FT DOMAIN 1 70 SER-RICH.
FT DOMAIN 117 137 SER-RICH.
FT DOMAIN 272 364 CHROMO DOMAIN.
FT DOMAIN 389 452 CHROMO DOMAIN.
FT NP_BIND 506 513 ATP (POTENTIAL).
FT SITE 614 617 DEAH BOX.
FT DOMAIN 1628 1644 3 X 5 AA REPEATS OF H-S-D-H-R.
FT REPEAT 1628 1632 1.
FT REPEAT 1634 1638 2.
FT REPEAT 1640 1644 3.
SQ SEQUENCE 1709 AA; 196517 MW; EC7F932A CRC32;

Query Match 67.3%; Score 33; DB 1; Length 1709;
Best Local Similarity 77.8%; Pred. No. 69;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLRLRNER 10
:|||||
Db 795 KLLRLRER 803

RESULT 7

CHDI_MOUSE
ID CHDI_MOUSE STANDARD; PRT; 1711 AA.
AC P40201;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE CHROMODOMAIN-HELICASE-DNA-BINDING PROTEIN 1 (CHD-1).
GN CHDI OR CHD-1.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 93211972.
RA DELMAS V., STOKES D.G., PERRY R.P.;
RT "A mammalian DNA-binding protein that contains a chromodomain and an

RT SNF2/SWI2-like helicase domain.";
 RL Proc. Natl. Acad. Sci. U.S.A. 90:2414-2418(1993).
 CC -!- FUNCTION: SEQUENCE-SELECTIVE DNA-BINDING PROTEIN. COULD PLAY AN
 CC IMPORTANT ROLE IN GENE REGULATION.
 CC -!- SUBCELLULAR LOCATION: NUCLEAR.
 CC -!- TISSUE SPECIFICITY: ABUNDANCE IS HIGHER IN CELLS REPRESENTING
 CC EARLY STAGES OF THE B LYMPHOID LINEAGE SUCH AS PRE-B AND B CELLS,
 CC THAN IN CELLS REPRESENTING MATURE PLASMACYTES OR OTHER CELL
 CC LINEAGES SUCH AS FIBROBLASTS.
 CC -!- SIMILARITY: BELONGS TO THE SNF2/RAD54 HELICASE FAMILY.
 CC -!- SIMILARITY: CONTAINS 2 'CHROMO' DOMAINS.
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 CC
 DR EMBL: L10410; AAB08486.1; -.
 DR PIR: A47392; A47392.
 DR HSP: P23197; LAP0.
 DR MGD: MGI:88393; CHD1.
 DR PROSITE: PS00598; CHROMO_1; 2.
 DR PROSITE: PS00013; CHROMO_2; 2.
 DR PFAM: PF00176; SNF2_N; 1.
 DR PFAM: PF00271; helicase_C; 1.
 DR PFAM: PF00385; chromo; 2.
 KW DNA-binding; ATP-binding; Helicase; Nuclear protein; Repeat.
 FT DOMAIN 1 70
 FT SER-RICH.
 FT DOMAIN 116 136
 FT SER-RICH.
 FT DOMAIN 270 362
 FT CHROMO DOMAIN.
 FT DOMAIN 387 450
 FT CHROMO DOMAIN.
 FT NP_BIND 504 511
 FT ATP (POTENTIAL).
 FT SITE 612 615
 FT DEAH BOX.
 FT DOMAIN 1629 1645
 FT 3 X 5 AA REPEATS OF H-S-D-H-R.
 FT REPEAT 1629 1633 1.
 FT REPEAT 1635 1639 2.
 FT REPEAT 1641 1645 3.
 FT REPEAT 1641 1645 3.
 SQ SEQUENCE 1711 AA; 196409 MW; CB184D33 CRC32;

Query Match 67.3%; Score 33; DB 1; Length 1711;
 Best Local Similarity 77.8%; Pred. No. 69;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLRLRLNER 10
 DB 793 KLLRLNER 801

RESULT 8
 Y487_MYCTU STANDARD; PRT; 183 AA.
 AC Q11153;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 15-DEC-1999 (Rel. 39, Last annotation update)
 DE HYPOTHETICAL 20.7 KD PROTEIN RV0487.
 GN RV0487 OR MTCY20G9.13.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-H37RV;
 RX MEDLINE: 98295987.
 RA COLE S.T., BROSCHE R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
 RA GORDON S.V., EIGLMEIER K., GAS S., BARRY C.E. III, TEKAIA F.,
 RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
 RA DAVIES R., DEVLIN K., FELTWELL T., GENTLES S., HAMLIN N., HOLROYD S.,
 RA HORNSBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,

RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAM M.A., ROGERS J.,
 RA RUTTER S., SEEGER K., SKELTON S., SQUARES S., SQUARES R., SULLSTON J.E.,
 RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 CC complete genome sequence.";
 RL Nature 393:537-544(1998).
 CC -!- SIMILARITY: STRONG, TO M.LEPRAE U2168E.
 CC
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 CC
 DR EMBL: Z77162; CAB00948.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 183 AA; 20716 MW; 23C141D2 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 183;
 Best Local Similarity 60.0%; Pred. No. 9.7;
 Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLRLNER 10
 DB 77 YRLRLNER 86

RESULT 9
 YF8_YEAST STANDARD; PRT; 221 AA.
 ID YF8_YEAST
 AC P43592;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 01-NOV-1995 (Rel. 32, Last annotation update)
 DE HYPOTHETICAL 25.9 KD PROTEIN IN MPRI-GCN20 INTERGENIC REGION.
 GN YFR008W.
 OS Saccharomyces cerevisiae (Baker's yeast).
 CC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 CC Saccharomycetaceae; Saccharomyces.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-S288C / AB972;
 RX MEDLINE: 95400292.
 RA MURAKAMI Y., NAITOU M., HAGIWARA H., SHIBATA T., OZAWA M.,
 RA SASANUMA S.-I., SASANUMA M., TSUCHIYA Y., SOEDA E., YOKOYAMA K.,
 RA YAMAZAKI M., TASHIRO H., EKI T.;
 RT "Analysis of the nucleotide sequence of chromosome VI from
 CC Saccharomyces cerevisiae.";
 RL Nat. Genet. 10:261-268(1995).
 CC
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 CC

Query Match 65.3%; Score 32; DB 1; Length 221;
 Best Local Similarity 75.0%; Pred. No. 12;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLRLNER 8
 DB 11111111


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Db 211 YRLRLRLH 218
RESULT 10
XJ9D_YEAST STANDARD; PRT; 239 AA.
ID YJ9D_YEAST
AC P47167;
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 01-OCT-1996 (Rel. 34, Last annotation update)
DE HYPOTHEICAL 27.6 KD PROTEIN IN NMD5-HOM6 INTERGENIC REGION.
GN YJR135C OR J2122.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomycetes.
RN [1]
RP SEQUENCE FROM N.A.
RA ROSE M., KOTTER P., ENTIAN K.D.;
RL Submitted (SEP-1995) to the EMBL/GenBank/DBJ databases.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: J04184; AAA34578.1; -
CC EMBL: U25842; AAB68116.1; -
CC PIR: A32122; A32122.
CC SGD: L0000524; DPM1.
CC PFAM: PF00535; Glycosyltransferase; Transmembrane;
KW Transferase; Glycosyltransferase; Transmembrane;
KW Endoplasmic reticulum.
FT TRANSMEM 239 259 POTENTIAL.
SQ SEQUENCE 239 AA; 27567 MW; 0BF23C5E CRC32;

Query Match 65.3%; Score 32; DB 1; Length 239;
Best Local Similarity 75.0%; Pred. No. 13;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRLN 8
|:|:|:|
Db 205 YRLRLRLN 212

RESULT 11
DPM1_YEAST STANDARD; PRT; 267 AA.
ID DPM1_YEAST
AC P14020;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE DOLICHOH-PHOSPHATE MANNOsylTRANSFERASE (EC 2.4.1.83) (DOLICHOH-
DE PHOSPHATE MANNOSE SYNTHASE) (DOLICHYL-PHOSPHATE BETA-D-
DE MANNOsylTRANSFERASE).
GN DPM1 OR SED3 OR YPR183W OR P9705.3.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomycetes.
RN [1]
RP SEQUENCE FROM N.A.
RA MEDLINE; 89034276.
RA ORLEAN P., ALBRIGHT C., ROBBINS P.W.;
RT "Cloning and sequencing of the yeast gene for dolichol phosphate
RT mannose synthase, an essential protein.";
RL J. Biol. Chem. 263:17499-17507(1988).
RN [2]
RP SEQUENCE FROM N.A.
RA STRAIN-5288C / AB972;
RA JOHNSTON M., ANDREWS S., BRINKMAN R., COOPER J., DING H., DU Z.,
RA FAVELLO A., FULTON L., GATTUNG S., GRECO T., KIRSTEN J.,
RA KUCABA T., HALLSWORTH K., HAWKINS J., HILLIER L., JTER M.,
RA JOHNSTON D., JOHNSTON L., LANGSTON Y., LATREILLE P., LE T.,
RA MARDIS E., MENEZES S., MILLER N., NHAN M., PAULEY A., PELUSO D.,
RA RIFKEN L., RILES L., TAICH A., TREVASKIS E., VIGNATI D.,
RA WILCOX L., WOHLDMAN P., VAUDIN M., WILSON R., WATERSTON R.;

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RL Submitted (APR-1995) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: INVOLVED IN THE SYNTHESIS OF THE SUGAR DONOR DOL-P-MAN
CC WHICH IS REQUIRED IN THE SYNTHESIS OF N-LINKED AND O-LINKED
CC OLIGOSACCHARIDES AND FOR THAT OF GPI ANCHORS.
CC -!- CATALYTIC ACTIVITY: GDP-MANNOSE + DOLICHYL PHOSPHATE -> GDP +
CC DOLICHYL D-MANNOSYL PHOSPHATE.
CC -!- SUBCELLULAR LOCATION: IF THE N-TERMINUS IS A FUNCTIONAL SIGNAL
CC SEQUENCE, THE PROTEIN IS PREDICTED TO BE ORIENTED TOWARD THE LUMEN
CC OF THE ENDOPLASMIC RETICULUM WITH BOTH TERMINI SERVING AS ANCHORS.
CC THE LACK OF A SIGNAL SEQUENCE INDICATES THAT THE ENZYME FACES THE
CC CYTOPLASM AND IS ANCHORED AT THE C-TERMINUS.
CC -!- DOMAIN: THE N-TERMINUS OF THE PROTEIN, THOUGH NOT HYDROPHOBIC,
CC MEETS EXISTING CRITERIA FOR YEAST SIGNAL SEQUENCES, EVEN THOUGH NO
CC SITE EXISTS FOR CLEAVAGE BY SIGNAL PEPTIDASE.
CC -!- SIMILARITY: BELONGS TO THE GLYCOSYLTRANSFERASE FAMILY 2.
CC -----
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CC -----
CC EMBL: J04184; AAA34578.1; -
CC EMBL: U25842; AAB68116.1; -
CC PIR: A32122; A32122.
CC SGD: L0000524; DPM1.
CC PFAM: PF00535; Glycosyltransferase; Transmembrane;
KW Transferase; Glycosyltransferase; Transmembrane;
KW Endoplasmic reticulum.
FT TRANSMEM 239 259 POTENTIAL.
SQ SEQUENCE 267 AA; 30362 MW; F8D92784 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 267;
Best Local Similarity 55.6%; Pred. No. 15;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLLRLRLN 10
|:|:|:|
Db 65 RIIVRTNR 73

RESULT 12
RL5_ANOGA STANDARD; PRT; 327 AA.
ID RL5_ANOGA
AC Q44248;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE 60S RIBOSOMAL PROTEIN L5.
GN RPL5
OS Anopheles gambiae (African malaria mosquito).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Nematocera; Culicoidae;
OC Culicidae; Anophelinae.
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN-63;
RA CORNEL A.J., KUMAR V., MUKABAYIRE O., SALAZAR RAFFERTY C.,
RA PETRARCA V., COLUZZI M., COLLINS F.H.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: THIS PROTEIN BINDS 5S RNA (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE L18P FAMILY OF RIBOSOMAL PROTEINS.
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CC EMBL; AF002238; AAB97731.1; -
DR PFAM; PF00061; Ribosomal_L18p; 1.
KW Ribosomal protein; rRNA-binding.
SQ SEQUENCE 327 AA; 37996 MW; F3A3ED2 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 327;
Best Local Similarity 50.0%; Pred. No. 18;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLNER 10
DB 49 FRLVRLSNR 58

RESULT 13
SAPD_ECOLI STANDARD; PRT; 330 AA.
AC P36635;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE PEPTIDE TRANSPORT SYSTEM ATP-BINDING PROTEIN SAPD.
GN SAPD.
OS Escherichia coli.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-K12;
RA EPSTEIN W., NOELKER E., STUMPE S., TEWES R., SCHMID R., BAKKER E.P.;
RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-K12 / MG1655;
RX MEDLINE; 9742617.
RA BLATTNER F.R., PLUNKETT G. III, BLOCH C.A., PERNA N.T., BURLAND V.,
RA RILEY M., COLLADO-VIDES J., GLASNER F.D., RODE C.K., MAYHEW G.F.,
RA GREGOR J., DAVIS N.W., KIRKPATRICK H.A., GOEDEN M.A., ROSE D.J.,
RA MAU B., SHAO Y.;
RT "The complete genome sequence of Escherichia coli K-12.";
RL Science 277:1453-1474(1997).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-K12;
RX MEDLINE; 97251357.
RA AIBA H., BABA T., FUJITA K., HAYASHI K., INADA T., ISONO K.,
RA ITOH T., KASAI H., KASHIMOTO K., KIMURA S., KITAKAWA M.,
RA KITAGAWA M., MAKINO K., MIKI T., MIZOBUCHI K., MORI H., MORI T.,
RA MOTOMURA K., NAKADE S., NAKAMURA Y., NASHIMOTO H., NISHIO Y.,
RA OSHIMA T., SAITO N., SAMPEI G., SEKI Y., SIVASUNDARAM S.,
RA TAGAMI H., TAKEDA J., TAKEMOTO K., TAKEUCHI Y., WADA C.,
RA YANAMOTO Y., HORIUCHI T.;
RT "A 570-kb DNA sequence of the Escherichia coli K-12 genome
corresponding to the 28.0-40.1 min region on the linkage map.";
RL DNA Res. 3:363-377(1996).
RN [4]
RP SEQUENCE OF 301-330 FROM N.A.
RA BERGLER H., EBELING A., FUCHSICHLEH S., HOGENAUER G.,
RA TURNOWSKY F.;
RL Submitted (MAR-1994) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: INVOLVED IN A PEPTIDE INTAKE TRANSPORT SYSTEM THAT
PLAYS A ROLE IN THE RESISTANCE TO ANTIMICROBIAL PEPTIDES.
CC -!- SUBCELLULAR LOCATION: INNER MEMBRANE-ASSOCIATED (POTENTIAL).
CC -!- SIMILARITY: BELONGS TO THE ATP-BINDING TRANSPORT PROTEIN FAMILY
(ABC TRANSPORTERS).
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CC
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CC
CC EMBL; X97282; CAA65940.1; -
DR EMBL; AE000227; AAC74373.1; -
DR EMBL; D90766; CAB20760.1; -
DR EMBL; D90767; CAB20768.1; -
DR EMBL; D90768; CAB20789.1; -
DR EMBL; U08190; AAA17670.1; -
DR EMBL; EG12304; SAPD.
DR PROSITE; PS00211; ABC_TRANSPORTER; FALSE_NEG.
DR PFAM; PF00005; ABC_tran; 1.
KW Peptide transport; Transport; Inner membrane; ATP-binding.
FT NP_BIND 40 47 ATP (POTENTIAL).
SQ SEQUENCE 330 AA; 37660 MW; BE092E6 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 330;
Best Local Similarity 66.7%; Pred. No. 18;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLNER 9
DB 198 FRLRLNQ 206

RESULT 14
SAPD_SALTY STANDARD; PRT; 330 AA.
AC P36636;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE PEPTIDE TRANSPORT SYSTEM ATP-BINDING PROTEIN SAPD.
GN SAPD.
OS Salmonella typhimurium.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Salmonella.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-ATCC 14028S;
RX MEDLINE; 94038887.
RA PARRA-LOPEZ C., BAER M.T., GROISMAN E.A.;
RT "Molecular genetic analysis of a locus required for resistance to
antimicrobial peptides in Salmonella typhimurium.";
RL EMBO J. 12:4053-4062(1993).
CC -!- FUNCTION: INVOLVED IN A PEPTIDE INTAKE TRANSPORT SYSTEM THAT
PLAYS A ROLE IN THE RESISTANCE TO ANTIMICROBIAL PEPTIDES.
CC -!- SUBCELLULAR LOCATION: INNER MEMBRANE-ASSOCIATED (POTENTIAL).
CC -!- SIMILARITY: BELONGS TO THE ATP-BINDING TRANSPORT PROTEIN FAMILY
(ABC TRANSPORTERS).
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or send an email to license@isb-sib.ch).
CC
CC EMBL; X74212; CAA52287.1; -
DR PIR; S39588; S39588.
DR SYGENE; SG10380; SAPD.
DR PROSITE; PS00211; ABC_TRANSPORTER; FALSE_NEG.
DR PFAM; PF00005; ABC_tran; 1.
KW Peptide transport; Transport; Inner membrane; ATP-binding.
FT NP_BIND 40 47 ATP (POTENTIAL).
SQ SEQUENCE 330 AA; 37611 MW; 9910EB90 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 330;
Best Local Similarity 66.7%; Pred. No. 18;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;
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Oy 1 YRLIRLNE 9
Db 198 FRLRLNQ 206

RESULT 15
VGLC_HSVB STANDARD; PRT; 468 AA.
AC P12889; P36321;
DT 01-OCT-1989 (Rel. 12, Created)
DT 01-OCT-1989 (Rel. 12, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE GLYCOPROTEIN C PRECURSOR (GLYCOPROTEIN 13).
GC OR GP13 OR 16.
OS Equine herpesvirus type 1 (strain Ab4p) (EHV-1), and
OS Equine herpesvirus type 1 (strain Kentucky D) (EHV-1).
OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
OC Alphaherpesvirinae; Varicellovirus.
RN [1]
RN SEQUENCE FROM N.A.
RP STRAIN=AB4P;
RX MEDLINE: 92295566.
RA TELFORD E.A.R., WATSON M.S., MCBRIDE K., DAVISON A.J.;
RT "the DNA sequence of equine herpesvirus-1.";
RN [2]
RN SEQUENCE FROM N.A.
RP STRAIN=KENTUCKY D;
RX MEDLINE: 88275055.
RA ALLEN G.P., COOGLER L.D.;
RT "Characterization of an equine herpesvirus type 1 gene encoding a
RT glycoprotein (gp13) with homology to herpes simplex virus
RT glycoprotein C.";
RL J. Virol. 62:2850-2858(1988).
RN [3]
RN SEQUENCE FROM N.A.
RP STRAIN=KENTUCKY D;
RX MEDLINE: 89382761.
RA GUO P., GOEBEL S., DAVIS S., PERKUS M.E., LANGUET B., DESMETTRE P.,
RA ALLEN G., PAOLETTI E.;
RT "Expression in recombinant vaccinia virus of the equine herpesvirus 1
RT gene encoding glycoprotein gp13 and protection of immunized
RT animals.";
RL J. Virol. 63:4189-4198(1989).
RN [4]
RN SEQUENCE FROM N.A.
RP STRAIN=KENTUCKY D;
RX MEDLINE: 93212524.
RA MATSUMURA T., SMITH R.H., O'CALLAGHAN D.J.;
RT "DNA sequence and transcriptional analyses of the region of the
RT equine herpesvirus type 1 Kentucky A strain genome encoding
RT glycoprotein C.";
RL Virology 193:910-923(1993).
CC -!- SIMILARITY: TO OTHER HERPESVIRUSES GLYCOPROTEIN C.
CC -!- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN GENE SUPERFAMILY.
CC -----
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CC -----
DR EMBL: L07272; AAA46078.1; -
DR EMBL: M86664; AAB02451.1; -
DR EMBL: M19966; AAA46077.1; -
DR EMBL: M29234; AAA46085.1; -
DR EMBL: S57839; AAB25944.1; -
DR PIR: H36796; VGBEAL.
DR PIR: A28149; VGBEEH.
DR PIR: B46114; B46114.

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KW Glycoprotein; Transmembrane; Signal.
FT SIGNAL 1 30 POTENTIAL.
FT CHAIN 31 468 GLYCOPROTEIN C.
FT TRANSMEM 432 451 POTENTIAL.
FT CARBOHYD 46 46 POTENTIAL.
FT CARBOHYD 57 57 POTENTIAL.
FT CARBOHYD 62 62 POTENTIAL.
FT CARBOHYD 92 92 POTENTIAL.
FT CARBOHYD 100 100 POTENTIAL.
FT CARBOHYD 131 131 POTENTIAL.
FT CARBOHYD 203 203 POTENTIAL.
FT CARBOHYD 208 208 POTENTIAL.
FT CARBOHYD 269 269 POTENTIAL.
FT CONFLICT 107 107 E -> K (IN REF. 4).
FT CONFLICT 145 145 E -> K (IN REF. 4).
FT CONFLICT 275 275 V -> A (IN REF. 4).
SQ SEQUENCE 468 AA; 50889 MW; EBF20B67 CRC32;

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Query Match 65.3%; Score 32; DB 1; Length 468;
Best Local Similarity 70.0%; Pred. No. 27;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

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Oy 1 YRLIRLNE 10
Db 104 YRLEIYNQR 113

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Search completed: February 8, 2000, 00:59:52
Job time: 3781 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:37 ; Search time 209.03 Seconds
(without alignments)
3.317 Million cell updates/sec

Title: US-08-653-294-11
Perfect score: 49
Sequence: 1 YRLIRLNR 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 225878 seqs, 69334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : SPITREMBL_12.*

- 1: sp-archaea.*
- 2: sp-bacteria.*
- 3: sp-fungi.*
- 4: sp-human.*
- 5: sp-invertebrate.*
- 6: sp-mammal.*
- 7: sp-mhc.*
- 8: sp-organelle.*
- 9: sp-phage.*
- 10: sp-plant.*
- 11: sp-rodent.*
- 12: sp-virus.*
- 13: sp-vertebrate.*
- 14: sp-unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	36	73.5	395	10 Q9ZT98	Q9ZT98 arabidopsis
2	34	69.4	254	11 P70561	P70561 rattus norv
3	34	69.4	485	12 Q39258	Q39258 equine herp
4	33	67.3	191	12 Q98543	Q98543 paramecium
5	33	67.3	201	3 Q12530	Q12530 saccharomyc
6	33	67.3	506	11 Q62157	Q62157 mus musculu
7	33	67.3	515	2 Q9ZAP8	Q9ZAP8 paenibacilli
8	33	67.3	537	5 Q22520	Q22520 caenorhabdi
9	33	67.3	560	13 Q08781	Q08781 gallus gall
10	33	67.3	859	11 P70261	P70261 mus musculu
11	33	67.3	1808	13 Q42142	Q42142 gallus gall
12	32	65.3	151	1 O58407	O58407 pyrococcus
13	32	65.3	261	5 Q96646	Q96646 drosophila
14	32	65.3	678	2 P71707	P71707 mycobacteri
15	31	63.3	214	10 Q65654	Q65654 arabidopsis
16	31	63.3	260	5 Q76669	Q76669 caenorhabdi
17	31	63.3	329	2 Q86190	Q86190 erwinia chr
18	31	63.3	397	10 Q48758	Q48758 arabidopsis
19	31	63.3	418	2 Q9X8L1	Q9X8L1 streptomyce
20	31	63.3	425	5 Q19125	Q19125 caenorhabdi

21	31	63.3	442	8 Q9XMT9	Q9XMT9 tetrahymena
22	31	63.3	570	3 Q13347	Q13347 magnaporthe
23	31	63.3	688	8 Q34312	Q34312 dictyosteli
24	31	63.3	819	3 Q94254	Q94254 schizosacch
25	31	63.3	1097	3 Q13592	Q13592 saccharomyc
26	31	63.3	1131	10 Q23741	Q23741 brassica ol
27	30	61.2	81	12 Q9WI79	Q9WI79 human immun
28	30	61.2	144	5 Q27369	Q27369 trypanosoma
29	30	61.2	173	2 Q51316	Q51316 nostoc sp.
30	30	61.2	175	5 Q17265	Q17265 brugia paha
31	30	61.2	176	2 Q32098	Q32098 bacillus su
32	30	61.2	186	2 Q67378	Q67378 aquifex aeo
33	30	61.2	189	12 Q98542	Q98542 paramecium
34	30	61.2	216	2 Q85823	Q85823 versinia ps
35	30	61.2	296	5 Q9Y0H6	Q9Y0H6 myxine glut
36	30	61.2	309	13 Q91634	Q91634 xenopus lae
37	30	61.2	366	2 Q9ZBT9	Q9ZBT9 streptomyce
38	30	61.2	367	10 Q80931	Q80931 arabidopsis
39	30	61.2	378	2 Q87101	Q87101 bacillus su
40	30	61.2	394	8 Q34835	Q34835 kluyveromyc
41	30	61.2	431	13 Q9W734	Q9W734 gallus gall
42	30	61.2	437	5 Q19360	Q19360 caenorhabdi
43	30	61.2	447	5 Q21147	Q21147 caenorhabdi
44	30	61.2	460	2 P94248	P94248 bifidobacte
45	30	61.2	471	5 Q9Y1X6	Q9Y1X6 ephydatia f

ALIGNMENTS

RESULT 1

Q9ZT98
ID Q9ZT98 PRELIMINARY; PRT; 395 AA.
AC Q9ZT98;
DT 01-MAY-1999 (TRENBLrel. 10, Created)
DT 01-MAY-1999 (TRENBLrel. 10, Last sequence update)
DE 01-MAY-1999 (TRENBLrel. 10, Last annotation update)
DE PUTATIVE LEUCINE-RICH REPEAT PROTEIN.
OS T419.11.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC eubryophytes; Spermatophyta; Magnoliophyta; eudicotyledons;
OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
OC Arabidopsis.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-CV. COLUMBIA;
RA PARNELL L.D., GNOJ L., DE LA BASTIDE M., HAMEED A., HABERMANN K.,
RA SCHUTZ K., HUANG E., GOTTESMAN T., DEDHIA N.N., MCCOMBIE W.R.;
RT "Genomic sequence of BAC T419 from Arabidopsis thaliana, Chromosome
IV, near 16.6 CM.";
RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF069442; AAC79105.1;
SQ SEQUENCE 395 AA; 43507 MW; F547BBD5 CRC32;

Query Match 73.5%; Score 36; DB 10; Length 395;

Best Local Similarity 66.7%; Pred. No. 15;

Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YRLIRLNR 9

Db 231 YRVLLRLNQ 239

RESULT 2

P70561
ID P70561 PRELIMINARY; PRT; 254 AA.
AC P70561;
DT 01-FEB-1997 (TRENBLrel. 02, Created)
DT 01-FEB-1997 (TRENBLrel. 02, Last sequence update)
DT 01-AUG-1998 (TRENBLrel. 07, Last annotation update)
DE FGF RECEPTOR ACTIVATING PROTEIN FRAG1.

GN FRAG1.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=BRAIN;
 RA LORENZI M.V., HORII Y., YAMANAKA R., SAKAGUCHI K., MIKI T.;
 RL Proc. Natl. Acad. Sci. U.S.A. 0:0-0(0).
 DR EMBL: U57715; AAB07050.1; -;
 SQ SEQUENCE 254 AA; 23395 MW; 8470603F CRC32;

Query Match 69.4%; Score 34; DB 11; Length 254;
 Best Local Similarity 87.5%; Pred. No. 25;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLN 8
 |||||
 DB 109 YRLCRLN 116

RESULT 3
 ID O39258 PRELIMINARY; PRT; 485 AA.
 AC O39258;
 DT 01-JAN-1998 (TReMBLrel. 05, Created)
 DT 01-JAN-1998 (TReMBLrel. 05, Last sequence update)
 DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
 DE COUNTERPART OF HSV-1 GENE UL44 AND VZV GENE 14.
 GN 16.
 OS Equine herpesvirus 4.
 OC Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
 OC Alphaherpesvirinae; Varicellovirus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NS80567;
 RX MEDLINE; 91021040.
 RA NICOLSON L., ONIONS D.E.;
 RT "The nucleotide sequence of the equine herpesvirus 4 gC gene
 homologues";
 RL Virology 179:378-387(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NS80567;
 RX MEDLINE; 98264497.
 RA TELFORD E.A.R., WATSON M.S., PERRY J., CULLINANE A.A., DAVIDSON A.J.;
 RT "The DNA sequence of equine herpesvirus-4";
 RL J. Gen. Virol. 79:1197-1203(1998).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NS80567;
 RA TELFORD E.A., WATSON M.S., PERRY J., CULLINANE A.A., DAVIDSON A.J.;
 RL Submitted (OCT-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF030027; AAC59330.1; -;
 DR PRINTS; PR00668; GLYCOPROTEIN.
 SQ SEQUENCE 485 AA; 52539 MW; 8DF52A42 CRC32;

Query Match 69.4%; Score 34; DB 12; Length 485;
 Best Local Similarity 70.0%; Pred. No. 48;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLIRLNER 10
 |||||
 DB 120 YRLHLNQR 129

RESULT 4
 ID Q98543 PRELIMINARY; PRT; 191 AA.
 AC Q98543;
 DT -01-FEB-1997 (TReMBLrel. 02, Created)

DT 01-FEB-1997 (TReMBLrel. 02, Last sequence update)
 DT 01-NOV-1998 (TReMBLrel. 08, Last annotation update)
 DE GENOME, PARTIAL SEQUENCE.
 GN A4931.
 OS Paramesitium bursaria chlorella virus 1 (PBCV-1).
 OC Viruses; dsDNA viruses, no RNA stage; Phycodnaviridae; Phycodnavirus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 96400190.
 RA KUTISH G.F., LI Y., LU Z., FURUTA M., ROCK D.L., VAN ETEN J.L.;
 RT "Analysis of 76 kb of the chlorella virus PBCV-1 330-kb genome: map
 positions 182 to 258";
 RL Virology 223:303-317(1996).
 DR EMBL: U42580; AAC96860.1; -;
 SQ SEQUENCE 191 AA; 22651 MW; E4547C83 CRC32;

Query Match 67.3%; Score 33; DB 12; Length 191;
 Best Local Similarity 66.7%; Pred. No. 30;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLNE 9
 |||||
 DB 74 YRLVVRENE 82

RESULT 5
 ID Q12530 PRELIMINARY; PRT; 201 AA.
 AC Q12530;
 DT 01-NOV-1996 (TReMBLrel. 01, Created)
 DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
 DT 01-NOV-1996 (TReMBLrel. 01, Last annotation update)
 DE CHROMOSOME XII READING FRAME ORF YLR145W.
 GN 19634.3.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Saccharomyces.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA RIEGER M., MUELLER-AUER S., BRUECKNER M.;
 RL Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA MIPS;
 RL Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C (AB972);
 RA FULTON L.;
 RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C (AB972);
 RA JOHNSTON M., ANDREWS S., BRINKMAN R., COOPER J., DING H., DU Z.,
 RA FAVELLO A., FULTON L., GATTUNG S., GRECO T., KIRSTEN J., KUCABA T.,
 RA HALLSWORTH K., HAWKINS J., HILLIER L., JIER M., JOHNSON D.,
 RA JOHNSTON L., LANGSTON Y., LATREILLE P., LE T., MARDIS E., MENEZES S.,
 RA MILLER N., NHAN M., PAULEY A., PELUSO D., RIFKEN L., RILES L.,
 RA TAICH A., TREVASKIS E., VIGNATI D., WILCOX L., WOHLDMAN P., VAUDIN M.,
 RA WILSON R., WATERSTON R.;
 RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C (AB972);
 RA WATERSTON R.;
 RL Submitted (APR-1996) to the EMBL/GenBank/DBJ databases.
 DR EMBL: Z73317; CAA97717.1; -;
 DR EMBL: U53879; AAB82379.1; -;
 SQ SEQUENCE 201 AA; 23618 MW; FDD081D6 CRC32;

Query Match 67.3%; Score 33; DB 3; Length 201;

Best Local Similarity 60.08; Pred. No. 32;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLRLNER 10
| | | | |
Db 15 YRLILLNHR 24

RESULT 6
ID Q62157 PRELIMINARY; PRT; 506 AA.
AC Q62157;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE ZINC FINGER PROTEIN (FRAGMENT).
GN RFP.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=BALE/C;
RA TAKAHASHI M.;
RL Submitted (OCT-1993) to the EMBL/GenBank/DBJ databases.
CC -1- SIMILARITY: CONTAINS A C3HC4-CLASS ZINC FINGER.
DR EMBL; X75343; CAA53092.1; -;
DR PROSITE; PS00518; ZINC_FINGER_C3HC4; 1.
DR PFAM; PF00622; SPRV; 1.
DR PFAM; PF00643; zf-B-box; 1.
DR PFAM; PF00097; zf-C3HC4; 1.
DR PFAM; PF00097; zf-C3HC4; 1.
KW DNA-binding; Zinc-finger.
FT NON_TER 1
SQ SEQUENCE 506 AA; 57882 MW; AEE397C3 CRC32;

Query Match 67.38; Score 33; DB 11; Length 506;
Best Local Similarity 77.88; Pred. No. 79;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLRLNER 9
| | | | |
Db 191 YRLRLNER 199

RESULT 7
ID Q9ZAP8 PRELIMINARY; PRT; 515 AA.
AC Q9ZAP8;
DT 01-MAY-1999 (TReMBLrel. 10, Created)
DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE NEOPULLULANASE.
OS Paenibacillus polymyxa (Bacillus polymyxa).
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Paenibacillus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CECT 155;
RX MEDLINE; 99118304.
RA YEBRA M.J., BLASCO A., SANZ P.;
RT "Expression and secretion of Bacillus polymyxa neopullulanase in
Saccharomyces cerevisiae.";
RL FEMS Microbiol. Lett. 170:41-49(1999).
DR EMBL; U89716; AAD05199.1; -;
DR HSSP; P21332; LUOK.
SQ SEQUENCE 515 AA; 58749 MW; 68075B21 CRC32;

Query Match 67.38; Score 33; DB 2; Length 515;
Best Local Similarity 77.88; Pred. No. 81;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLRLNER 9
| | | | |
Db 419 YRLRLNER 427

RESULT 8
ID Q22520 PRELIMINARY; PRT; 537 AA.
AC Q22520;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
DT 01-JAN-1999 (TReMBLrel. 09, Last annotation update)
DE T16A9.1 PROTEIN.
GN T16A9.1.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RA MCMURRAY A.;
RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA WILSON R., AINSOUGH R., ANDERSON K., BAYNES C., BERKS M.,
RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
RA LIGHNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans.";
RL Nature 368:32-38(1994).
DR EMBL; Z77135; CAB00875.1; -;
SQ SEQUENCE 537 AA; 61447 MW; A7597F26 CRC32;

Query Match 67.38; Score 33; DB 5; Length 537;
Best Local Similarity 55.88; Pred. No. 84;
Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLRLNER 10
| | | | |
Db 113 KLIVKNER 121

RESULT 9
ID Q08781 PRELIMINARY; PRT; 560 AA.
AC Q08781;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
DT 01-MAY-1999 (TReMBLrel. 10, Last annotation update)
DE HYPOTHETICAL 64.4 KD PROTEIN.
OS Gallus gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Archosauria; Aves;
OC Neognathae; Galliformes; Phasianidae; Phasianinae; Gallus.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=LENS;
RX MEDLINE; 94116444.
RA FUNAHASHI J., SEKIDO R., MURAI K., KAWACHI Y., KONDOH H.;
RT "Delta-crystallin enhancer binding protein delta EFL is a zinc finger-
homeodomain protein implicated in postgastrulation embryogenesis.";
RL Development 119:433-446(1993).
DR EMBL; D14316; BAA03262.1; -;
DR PFAM; PF00271; helicase_C; 1.
KW Hypothetical protein.
SQ SEQUENCE 560 AA; 64394 MW; A54C9E16 CRC32;

Query Match 67.3%; Score 33; DB 13; Length 560;
 Best Local Similarity 77.8%; Pred. No. 88;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLLRLRNER 10
 :||||| ||
 Db 43 KLLRLRER 51

RESULT 10
 P70261 PRELIMINARY; PRT; 859 AA.
 AC P70261;
 DT 01-FEB-1997 (TREMELrel. 02, Created)
 DT 01-FEB-1997 (TREMELrel. 02, Last sequence update)
 DT 01-AUG-1998 (TREMELrel. 07, Last annotation update)
 DE PALADIN GENE.
 GN PALADIN.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA PEARCE J.J.H., DAVIES T., GARDENER R.L.;
 RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
 DR EMBL; X99384; CAA67763.1; -.
 SQ SEQUENCE 859 AA; 96739 MW; 8D061D00 CRC32;

Query Match 67.3%; Score 33; DB 11; Length 859;
 Best Local Similarity 75.0%; Pred. No. 1.3e+02;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRNER 8
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 Db 464 YRLRLVNL 471

RESULT 11
 O42142 PRELIMINARY; PRT; 1808 AA.
 ID O42142;
 AC O42142;
 DT 01-JAN-1998 (TREMELrel. 05, Created)
 DT 01-JAN-1998 (TREMELrel. 05, Last sequence update)
 DT 01-NOV-1999 (TREMELrel. 12, Last annotation update)
 DE CHROMO-HELICASE-DNA-BINDING ON THE Z CHROMOSOME PROTEIN.
 GN CHD-2.
 OS Gallus gallus (Chicken).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Archosauria; Aves;
 OC Neognathae; Galliformes; Phasianidae; Phasianinae; Gallus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 97473516.
 RA GRIFFITHS R., KORN R.M.;
 RT "A CHD1 gene is z chromosome linked in the chicken Gallus
 domesticus";
 RN Gene 197:225-229(1997).
 DR EMBL; AF004397; AAC60282.1; -.
 DR HSSP; P23197; IAP0.
 DR PROSITE; PS00598; CHROMO_1; 2.
 DR PFAM; PF00385; chromo; 2.
 DR PFAM; PF00271; helicase_C; 1.
 DR PFAM; PF00176; SNE2_N; 1.
 KW Helicase; DNA-binding.
 SQ SEQUENCE 1808 AA; 208399 MW; 97FE8926 CRC32;

Query Match 67.3%; Score 33; DB 13; Length 1808;
 Best Local Similarity 77.8%; Pred. No. 2.8e+02;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLLRLRNER 10
 :||||| ||
 Db 792 KLLRLRER 800

RESULT 12
 O38407 PRELIMINARY; PRT; 151 AA.
 ID O38407;
 AC O58407;
 DT 01-AUG-1998 (TREMELrel. 07, Created)
 DT 01-AUG-1998 (TREMELrel. 07, Last sequence update)
 DT 01-JAN-1999 (TREMELrel. 09, Last annotation update)
 DE 151AA LONG HYPOTHETICAL FRXA PROTEIN.
 GN PH0674.
 OS Pyrococcus horikoshii.
 OC Archaea; Euryarchaeota; Thermococcales; Thermococcaceae; Pyrococcus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-OT3;
 RX MEDLINE; 98344137.
 RA KAWARABAYASI Y., SAWADA M., HORIKAWA H., HAIKAWA Y., HINO Y.,
 RA YAMAMOTO S., SEKINE M., BABA S., KOSUGI H., HOSOIYAMA A., NAGAI Y.,
 RA SAKAI M., OGURA K., OTUKA R., NAKAZAWA H., TAKAMIYA M., OHFUKU Y.,
 RA FUNAHASHI T., TANAKA T., KUDOH Y., YAMAZAKI J., KUSHIDA N., OGUCHI A.,
 RA AOKI K., NAKAMURA Y., ROBB T.F., HORIKOSHI K., MASUCHI Y., SHIZUYA H.,
 RA KIKUCHI H.;
 RT "Complete sequence and gene organization of the genome of a hyper-
 thermophilic archaeobacterium, Pyrococcus horikoshii OT3";
 RL DNA Res. 5:55-76(1998).
 DR EMBL; AP000003; BAA29765.1; -.
 SQ SEQUENCE 151 AA; 17160 MW; 11AACD59 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 151;
 Best Local Similarity 77.8%; Pred. No. 39;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLLRLRNER 10
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 Db 96 RLLIELDER 104

RESULT 13
 O96646 PRELIMINARY; PRT; 261 AA.
 ID O96646;
 AC O96646;
 DT 01-MAY-1999 (TREMELrel. 10, Created)
 DT 01-MAY-1999 (TREMELrel. 10, Last sequence update)
 DT 01-MAY-1999 (TREMELrel. 10, Last annotation update)
 DE PIGMENT CELL DEHYDROGENASE REDUCTASE.
 GN PCDR.
 OS Drosophila melanogaster (Fruit fly).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
 OC Ephydroidea; Drosophilidae; Drosophila.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 98384483.
 RA BRUNEL C.A., MADIGAN S.J., CASSILL J.A., EDEEN P.T., MCKEOWN M.;
 RT "pcdr, a novel gene with sexually dimorphic expression in the pigment
 cells of the Drosophila eye";
 RL Dev. Genes Evol. 208:327-335(1998).
 DR EMBL; AF098864; AAC72391.1; -.
 SQ SEQUENCE 261 AA; 28302 MW; DC731F30 CRC32;

Query Match 65.3%; Score 32; DB 5; Length 261;
 Best Local Similarity 60.0%; Pred. No. 66;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRNER 10
 :||||| ||
 Db 207 YRLDELNKQ 216

RESULT 14

P71707 PRELIMINARY; PRT; 678 AA.
AC P71707;
DT 01-NOV-1998 (TEMBLrel. 08, Created)
DT 01-NOV-1998 (TEMBLrel. 08, Last sequence update)
DT 01-MAY-1999 (TEMBLrel. 10, Last annotation update)
DE PROBABLE PENICILLIN-BINDING PROTEINS 1A/1B (PBP1).
GN PONA OR RV0050 OR MTCV21.13.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacteriaceae; Mycobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RA COLE S.T., BROSC R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
RA GORDON S.V., EIGLMEIER K., GAS S., BARRY C.E., TEKAIA F., BADCOCK K.,
RA BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R., DAVIES R.,
RA DEVLIN K., FELTWELL T., GENTLES S., HAMLIN N., HOLROYD S., HORNSBY T.,
RA JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L., OLIVER S.,
RA OSBORNE J., QUAIL M.A., RAJANDREAM M.A., ROGERS J., RUTTER S.,
RA SEEGER K., SKELTON S., SQUARES S., SQUARES R., SULSTON J.E., TAYLOR K.,
RA WHITEHEAD S., BARRELL B.G.;
RL Nature 393:537-544(1998).
CC -!- FUNCTION: CELL WALL FORMATION.
CC -!- PATHWAY: FINAL STAGES IN PEPTIDOGLYCAN SYNTHESIS.
CC -!- SUBCELLULAR LOCATION: MEMBRANE ASSOCIATED (BY SIMILARITY).
CC -!- SIMILARITY: TO OTHER BACTERIAL CLASS 1A PENICILLIN-BINDING
CC PROTEINS.
DR EMBL; Z80775; CAB02529.1; -;
DR PFAM; PF00912; Transglycosyl; 1.
DR PFAM; PF00905; Transpeptidase; 1.
KW Peptidoglycan synthesis; Cell wall; Transmembrane.
FT ACT SITE 347 347 ACYLATED BY PENICILLIN (BY SIMILARITY).
FT DOMAIN 625 631 POLY-PRO.
FT DOMAIN 664 670 POLY-PRO.
SQ SEQUENCE 678 AA; 71150 MW; 58CC83F1 CRC32;

Query Match 65.3%; Score 32; DB 2; Length 678;
Best Local Similarity 62.5%; Pred. No. 1.7e+02;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLRLN 8
|||::||
Db 404 YRLMLKN 411

RESULT 15

O65654 PRELIMINARY; PRT; 214 AA.
AC O65654;
DT 01-AUG-1998 (TEMBLrel. 07, Created)
DT 01-AUG-1998 (TEMBLrel. 07, Last sequence update)
DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
DE HYPOTHETICAL 24.8 KD PROTEIN.
GN T19P19.60.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;
OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
OC Arabidopsis.
RN [1]
RP SEQUENCE FROM N.A.
RA BEVAN M., MONFORT A., CASACUBERTA E., PUIGDOMENECH P., HOHEISEL J.,
RA MEWES H.W., MAYER K.F.X., SCHUELLER C.;
RL Submitted (APR-1998) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA EU ARABIDOPSIS SEQUENCING PROJECT;
RL Submitted (APR-1998) to the EMBL/GenBank/DBJ databases.

DR EMBL; AL022605; CAA18753.1; -;
DR MENDEL; 29179; Arath;3410;29179.
KW Hypothetical protein.
SQ SEQUENCE 214 AA; 24813 MW; 110A2C72 CRC32;

Query Match 63.3%; Score 31; DB 10; Length 214;
Best Local Similarity 75.0%; Pred. NO. 87;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

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Search completed: February 8, 2000, 13:17:39
Job time: 32488 sec

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Date: Feb 8, 2000 4:38 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

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-MINMATCH=0.100 -LOPCL=0.000 -LOPEXT=0.000 -OGAPOP=4.500
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Search information block:

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Database sequences: 821193
Database length: 1518192014
Search time (sec): 11370.480000

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gb_in2:AC001780	+	40.00	103.29	991.28	! AC017780 Drosophila melanog
gb_htg3:AC009355	-	40.00	102.46	66958	! AC009355 Drosophila melanog
gb_pr3:AC005365	+	40.00	100.28	86130	! AC005365 Homo sapiens chrom
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gb_htg3:AC008327	-	40.00	93.66	185469	! AC008327 Drosophila melanog
gb_pr2:AC0017912	-	39.00	101.01	49261	! AC017912 Drosophila melanog
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gb_htg4:AC007054	-	39.00	90.22	171979	! AC007054 Drosophila melanog
gb_htg4:AC0006467	+	39.00	90.03	175695	! AC006467 Drosophila melanog
gb_htg3:AC008682	-	39.00	88.22	216649	! AC008682 Homo sapiens chrom
gb_ro:AB012933	-	38.00	122.79	81.19	! AB012933 Rattus norvegicus m
gb_p11:ATHSP881	+	38.00	117.48	160.57	! Y11839 A. thaliana hsp88.1 ge
gb_htg3:AC008847	+	38.00	98.65	40552	! AC008847 Homo sapiens chrom
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gb_htg4:AC010574	-	38.00	93.26	75163	! AC010574 Drosophila melanog
gb_htg7:AC017187	-	38.00	91.37	93588	! AC017187 Drosophila melanog
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gb_p12:ATAC007167	+	38.00	90.47	103874	! AC007167 Arabidopsis thalian
gb_htg3:AC009251	-	38.00	89.96	110262	! AC009251 Drosophila melanog
gb_htg4:AC010121	-	38.00	89.72	113253	! AC010121 Drosophila melanog
gb_htg4:AC010006	-	38.00	89.71	113508	! AC010006 Drosophila melanog
gb_htg3:AC010112	-	38.00	89.08	122073	! AC010112 Drosophila melanog
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gb_p12:AF142691	-	37.00	118.36	143.34	! AF142691 Pterocarpus indicus m
gb_p11:BNMTWAD3	+	37.00	118.28	144.90	! D13697 Brassica napus mitochon
gb_p12:AF142728	+	37.00	118.22	145.96	! AF142728 Robinia pseudocacia
gb_in1:CEG0762	-	37.00	100.29	20720	! Z32840 Caenorhabditis elegans
gb_htg5:AC0014301	+	37.00	96.98	2.2e+03	! AC0014301 Drosophila melanog
gb_p12:F19K23	-	37.00	86.92	8.1e+03	! AC000375 Sequence of BAC F19K
gb_htg2:AC007583	+	37.00	86.05	9.0e+03	! AC007583 Arabidopsis thalian
gb_pr3:HS3365019	+	37.00	82.81	1.4e+04	! AL096867 Human DNA sequenc
gb_htg4:AC011274	-	37.00	82.27	1.5e+04	! AC011274 Homo sapiens clone

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DEFINITION Oryza sativa genomic DNA, chromosome 1, clone:P0003H10.
ACCESSION AP000815
VERSION AP000815.1 GI:6498418
KEYWORDS
SOURCE Oryza sativa (cultivar:Nipponbare) DNA, clone:P0003H10.
ORGANISM Oryza sativa

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
Poaceae; Oryza.

REFERENCE 1 (bases 1 to 142418)

AUTHORS Sasaki,T., Matsumoto,T. and Yamamoto,K.

TITLE Oryza sativa nipponbare(GA3) genomic DNA, chromosome 1, PAC

clon:P0003H10

JOURNAL Published Only in DataBase (1999) In press

REFERENCE 2 (bases 1 to 142418)

AUTHORS Sasaki,T., Matsumoto,T. and Yamamoto,K.

TITLE Direct Submission

JOURNAL Submitted (30-NOV-1999) to the DDBJ/EMBL/GenBank databases. Takuji

Sasaki, National Institute of Agrobiological Resources, Rice Genome
Research Program; Kannondai 2-1-2, Tsukuba, Ibaraki 305-8602, Japan
(E-mail:tsasaki@abr.affrc.go.jp,
URL:http://www.dna.affrc.go.jp:82/, Tel:81-298-38-7441,
Fax:81-298-38-7466)

COMMENT The orientation of the sequence is from T7 to SP6 of the PAC clone.
Genes were predicted from the integrated results of the
following:GENSCAN1.0, BLASTN2.0, BLASTX2.0 as well as
SplicePredictor (October 1998 version). The genomic sequence was
searched against the non-redundant database NRP (PIR, SWISSPROT,
GENPEPT, PDB) from MAFF DNA bank and the cDNA sequence database at
RGP. Protein similarities of the coding regions were searched
against NRP with BLASTP2.0. ESTs represent the identified cDNA
sequences using BLASTN 2.0 with the corresponding DDBJ accession
no. and RGP clone ID.

Detailed information on assembly quality together with annotation
of this entry at http://www.dna.affrc.go.jp:82/genomicdata/GenomeFei
nished.html.

FEATURES

Location/Qualifiers

1. .142418

/organism="Oryza sativa"

/cultivar="Nipponbare"

/db_xref="taxon:4530"

/chromosome="1"

/clone="P0003H10"

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5069..5509)

/note="Similar to Arabidopsis thaliana DNA chromosome 4,

BAC clone F22K18 (AL035356)"

/protein_start=1

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/db_xref="GI:6498419"

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RHKDNQNSPLKSNAGPLSVLYITVTFVSVMFVGLTFLVLMSTNQTTFENFRY
VADKXENPNRGAISNIAEFCAGIPPSNNRNSWVAPPLEEDDDVSQOLPPRNGAD
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CDS

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10743..11006)
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A081294(E10057),A032455(S10086) correspond to a region
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ESGVDFAEFEQVLY"
CDS
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17726, 17860, 18400, 18587, 18681, 18749, 18826, 18916,
19028, 19162, 19323, 19379, 19349, 19441, 19450, 19509,
19523, 19585, 19589, 19701, 19800, 19941, 20057, 20294,
21399, 21586)
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AU032236(R3784), AU081351(E61905), AU031587(E61905),
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gene.; Similar to alien-like protein. (AC005623)"
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INDYVSGSASQNSLQEFYQNTLKALBEAKNERLWFTNLKCKINFDMGEYGRMSK
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PHRITMGITRECGGKHAERQWDAATDFEAFKNDYDEAGNPRRQICLKLYLVANML
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VLSMCMELNPEKDEQVQLLYSLIDNRIOGHIDQVKKLLEGRDEIRSHQVEYSAE
EHLPGVQWREYRAGCLGALGTCLHLVLMMLGFCPELNSANSPSETILIAS
ANRWFCHRTSKFWYGNVFRVYAPGTRTANTVPLPSSVANQSLSNQNEG
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CDS
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gene encoding calmodulin. (Z12828)"
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complement(join(24778, 24897, 25615, 25722, 25808, 25893,
26090, 26177, 26435, 26638, 27519, 27581, 27708, 27778,
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ALPQISLPYLMIRDLKVAKEQDIGYFASCVQAYASVCVKKEHQALGISL
VTSRAIALVWPAIGGLSOTPLHMHDDKEVIDALBAQDATSDLGTTGSESGSMR
GHTKSLKNWOLMSAITLYCVFSLHDYALFISLWAVSSRKYRGLSFTSODVGIVLA
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join(40347, 40358, 40468, 40680)
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IPYLNFFYIWWVCLSPFRYRPAKKICDFSVMKAMHGHR"
join(47081, 47084, 47194, 47313, 47469, 47584)
/notes="hypothetical protein"
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/translation="MSRGRAPATACGSVRYAAEVAPEDEDSARYALLLLRLRDS
AATAPPLARRIRDHGVPLLEDEDLLSVGPTLD"
join(48412, 48822, 49072, 49179, 49283, 49438, 49876, 50091,
50202, 50411, 50488, 50625, 50842, 50943, 51105, 51170,
51274, 51358, 51454, 51551)
/notes="Similar to Arabidopsis thaliana chromosome II BAC
T27A16 sequence; hypothetical protein. (AC005496)"
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RRRLRGTDVRSNTSSSSRRKRDHDDHGGYDGDGAGALLASVRRLLSGSAQDD
AAEGEAEDGQGPQKWAIVFLCFSAFLCNDRVNMSIATLPMFAEFGNPTVGL
IOSFFWGLLTQIAGGWADTVGGTVLGVFWISVATATLPAFAKGLGFLLVTR
AFMGVSGVAMPANNILSKWPVYSERSIALVSGMVLGSLVGLAFLPILNFGW
PSVYSGSLGVFWFSTWASKYSSPLEDPSIAEKKLITSQTTGGEYKIEPWGLI
LSRPFVWALIVSHFCHNWGFILLTWPTIYNQVKFNLETESGLFCVLPWLTMAVSAN
FGWADTLVSRGSLVTVVRKIMOSIGFLGPAFFLTQLSHIDSAMAVLNCACSGTD
AGSGLYSNHQDIPRYAGVILGSLNTAGVAGVFGTAATGYILOHGSWDDVFKVS
VLYVGLVWNLFGTEKIID"
complement(join(52616, 52685, 53228, 53299, 53395, 53466,
53657, 53800, 54663, 54734, 56026, 56234))
/notes="ESTs C97644(C60871), AU081285(C60871) correspond to
a region of the predicted gene.; Similar to L.esculentum
LRP gene. (X95269)"
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SWDPTLVNPTFWHTCDRAGVTRLDLGNLSGHLAELGHLEHLOYLEYKNNIQ
KITPAEGLSKNLISLDLYNNITGTPKELGKLSLVFLRLNDSNGPTPRDLAKI
SSLKVIDVSNNDLGGTIPSTGPFHEIPLNFDKNPRLEGPLOGLATYDNC"
59608, 59826
/notes="hypothetical protein"
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/protein_id="BAA87833.1"
/db_xref="GI:6498430"
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complement(join(63689, 63842, 63844, 64145))
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Quality: 41.00 Length: 9
Ratio: 4.556 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 88.889
alignment_block:
US-08-653-294-11 x AP000815/rev ..
Align seg 1/1 to reverse of: AP000815 from: 1 to: 142418
1 TyrArgLeuIleuIleArgLeuasnGlu 9
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8117 TACCGCTTGTAACTGTTGAACCA 8091
seq_name: gb_in1:CEC50B6

seq_documentation_block:
LOCUS CEC50B6 41322 bp DNA INV 02-SEP-1999
DEFINITION Caenorhabditis elegans cosmid C50B6, complete sequence.
ACCESSION Z81050
VERSION Z81050.1 GI:1627685
KEYWORDS HTG.
SOURCE
ORGANISM Caenorhabditis elegans.
Eukaryote; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
Rhabditina; Rhabditidae; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 41322)
Wilson, R., Ainscough, R., Anderson, K., Baynes, C., Berks, M.,
Bonfield, J., Burton, J., Connell, M., Copsey, T., Cooper, J.,
Coulson, A., Craxton, M., Dear, S., Du, Z., Durbin, R., Favello, A.,
Fulton, L., Gardner, A., Green, P., Hawkins, T., Hillier, L., Jier, M.,
Johnston, L., Jones, M., Kershaw, J., Kirsten, J., Laister, N.,
Latreille, P., Lightning, J., Lloyd, C., McMurray, A., Mortimore, B.,
O'Callaghan, M., Parsons, J., Percy, C., Rifkin, L., Roopra, A.,
Saunders, D., Showkhen, R., Smalton, N., Smith, A., Sonhammer, E.,
Staden, R., Sulston, J., Thierry-Mieg, J., Thomas, K., Vaudin, M.,
Vaughan, K., Waterston, R., Watson, A., Weinstock, L.,
Wilkinson-Sproat, J. and Wohlman, P.
2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans
JOURNAL Nature 368 (6466), 32-38 (1994)
MEDLINE 94150718
REFERENCE 2 (bases 1 to 41322)
AUTHORS Percy, C.
TITLE Direct Submission
JOURNAL Submitted (21-OCT-1996) Louis, MO 63110, USA. E-mail:
jes@sanger.ac.uk or rw@nemastode.wustl.edu
COMMENT Coding sequences below are predicted from computer analysis, using
Predictions from GeneFinder (P. Green, U. Washington), and other
available information.
For a graphical representation of this sequence and its analysis
see:
http://webace.sanger.ac.uk/cgi-
bin/displaydb-wormacsclass=Sequence&object=C50B6
Current sequence finishing criteria for the C. elegans genome
sequencing consortium are that all bases are either sequenced
unambiguously on both strands, or on a single strand with both a
dye primer and dye terminator reaction, from distinct subclones.
Exceptions are indicated by an explicit note.
IMPORTANT: This sequence is NOT necessarily the entire insert of
the specified clone. It may be shorter because we only sequence
overlapping sections once, or longer because we arrange for a small
overlap between neighbouring submissions.
IMPORTANT: This sequence is not the entire insert of clone C50B6.
It may be shorter because we only sequence overlapping sections
once, or longer because we arrange for a small overlap between
neighbouring submissions.
The true left end of clone C50B6 is at 1 in this sequence. The true
right end of clone C50B6 is at 2293 in
sequence Z81524.
The true left end of clone F32H5 is at 41219 in this sequence. The
start of this sequence (1..101) overlaps with the end of sequence
Z81040.
The end of this sequence (41219..41322) overlaps with the start of
sequence Z81524.
FEATURES
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/db_xref="taxon:6239"
/chromosome="v"
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EMBL:R02164 comes from this gene; cDNA EST CEMSA18R comes from this
gene; cDNA EST EMBL:D33193 comes from this gene; cDNA EST
EMBL:D35959 comes from this gene; cDNA EST EMBL:D17778
comes from this gene; cDNA EST EMBL:D72307 comes from this
gene; cDNA EST EMBL:D72719 comes from this gene; cDNA EST
EMBL:D74467 comes from this gene; cDNA EST EMBL:D75157
comes from this gene; cDNA EST EMBL:D75631 comes from this
gene; cDNA EST EMBL:D64450 comes from this gene; cDNA EST
EMBL:D64882 comes from this gene; cDNA EST EMBL:D65720
comes from this gene; cDNA EST EMBL:D65951 comes from this
gene; cDNA EST EMBL:D67440 comes from this gene; cDNA EST
EMBL:D68140 comes from this gene; cDNA EST EMBL:D69258
comes from this gene; cDNA EST EMBL:D69570 comes from this
gene; cDNA EST EMBL:C12197 comes from this gene; cDNA EST
EMBL:C10518 comes from this gene; cDNA EST Yk670d2.3 comes
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SVAASTAEVFGDAHEDTEANFLYKALLEGKLEDRVLNALTDPVPAEVEEVQ
DGIVENPDPQDEAEIKQVEEALGVAEEPEPTVADEAVKTEQEAEDSEKVEQ
NSDEQNEEEVVEATVDPTEDVEVEVEVEVEVEVEVEVEVEVEVEVEVEVEVE
EADQSMKLAWELLETSICADKKAASAEESTVDEGAIKMKLNADVLTSLGHGTAD
SKYEQAKDLTEAISIQTVHLPATSRVIANVTLLAKAFSSDSLSLFEQAAAHFNDTKNI
LIAKSELKQLETVSDDAKSDIQSEIKELDGLIPELDAFIVDARASAEOTEKLKSDA
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7495..7971,8025..8084)
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OLFTLKLAPGDTENLEEDENYAKLLFSCTITAVMVALHNDHEMSNSICPDYLKPEY
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/gene="C50B6.4"
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TITLE
JOURNAL
REFERENCE
AUTHORS

Rubin, G.M.
Sequencing of *Drosophila melanogaster*
2 (bases 1 to 66958)

Celniker, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazek, R.G.,
Butenkov, C., Champe, M., Chavez, C., Chew, M., Ciesiolka, L.,
Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L.,
Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L.,
Kim, E., Lee, B., Lewis, S., Li, P., Lomotan, M.A., Mazda, P.,
Moshrefi, A.R., Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S.,
Pfeiffer, B., Poon, L., Sequeira, A., Sethi, R., Snir, E.,
Svirskas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and
Rubin, G.M.

TITLE
JOURNAL

Submitted (17-AUG-1999), Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA

On Sep 20, 1999 this sequence version replaced gi:5748857.
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive web site (<http://www.fruitfly.org/sequence/>) or send email
to bdg@fruitfly.berkeley.edu. All contigs in this submission meet
the following cutoffs: length >= 200 bases.

* NOTE: This is a 'working draft' sequence. It currently
* consists of 76 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 1075: contig of 1075 bp in length

* 1076 1155: gap of unknown length
* 1156 1835: contig of 680 bp in length
* 1836 1915: gap of unknown length
* 1916 2905: contig of 990 bp in length
* 2906 2985: gap of unknown length
* 2986 3617: contig of 632 bp in length
* 3618 3697: gap of unknown length
* 3698 4682: contig of 985 bp in length
* 4683 4762: gap of unknown length
* 4763 5235: contig of 473 bp in length
* 5236 5315: gap of unknown length
* 5316 5956: contig of 641 bp in length
* 5957 6036: gap of unknown length
* 6037 6655: contig of 619 bp in length
* 6656 6735: gap of unknown length
* 6736 7335: contig of 600 bp in length
* 7336 7415: gap of unknown length
* 7416 8119: contig of 704 bp in length
* 8120 8199: gap of unknown length
* 8200 8769: contig of 570 bp in length
* 8770 8849: gap of unknown length
* 8850 9419: contig of 570 bp in length
* 9420 9499: gap of unknown length
* 9500 9915: contig of 416 bp in length
* 9916 9995: gap of unknown length
* 9996 11078: contig of 1083 bp in length
* 11079 11158: gap of unknown length
* 11159 12139: contig of 981 bp in length
* 12140 12219: gap of unknown length
* 12220 12951: contig of 732 bp in length
* 12952 13031: gap of unknown length
* 13032 13776: contig of 745 bp in length
* 13777 13856: gap of unknown length
* 13778 15280: contig of 1424 bp in length
* 13857 15281: gap of unknown length
* 15282 16485: contig of 1125 bp in length
* 16486 16565: gap of unknown length
* 16566 17573: contig of 1008 bp in length
* 17574 17653: gap of unknown length
* 17654 18313: contig of 660 bp in length
* 18314 18393: gap of unknown length
* 18394 19168: contig of 775 bp in length

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* 19249
* 20163
* 20243
* 21205
* 21285
* 22698
* 22778
* 24100
* 24180
* 24181
* 25936
* 25937
* 26017
* 27359
* 27639
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* 30696
* 30775
* 30776
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* 33680
* 33681
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* 36279
* 36359
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* 45180
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* 55185

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21284: gap of unknown length
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24100: contig of 1322 bp in length
24180: gap of unknown length
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26016: gap of unknown length
27358: contig of 1542 bp in length
27638: gap of unknown length
29229: contig of 1591 bp in length
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30695: contig of 1386 bp in length
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30776: contig of 1784 bp in length
32559: gap of unknown length
32639: gap of unknown length
33600: contig of 961 bp in length
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33681: contig of 1631 bp in length
35311: gap of unknown length
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36278: contig of 887 bp in length
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43592: contig of 555 bp in length
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45179: gap of unknown length
45825: contig of 646 bp in length
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46485: contig of 580 bp in length
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48534: contig of 622 bp in length
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49185: contig of 571 bp in length
49265: gap of unknown length
49805: contig of 540 bp in length
49885: gap of unknown length
50509: contig of 624 bp in length
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51193: contig of 604 bp in length
51273: gap of unknown length
51887: contig of 614 bp in length
51967: gap of unknown length
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52590: gap of unknown length
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53656: contig of 518 bp in length
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54452: contig of 716 bp in length
54532: gap of unknown length
55105: contig of 573 bp in length
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* 57799 58374: contig of 576 bp in length
* 58375 58454: gap of unknown length
* 58455 58951: contig of 497 bp in length
* 58952 59031: gap of unknown length
* 59032 59665: contig of 634 bp in length
* 59666 59745: gap of unknown length
* 59746 60315: contig of 570 bp in length
* 60316 60395: gap of unknown length
* 60396 60964: contig of 569 bp in length
* 60965 61044: gap of unknown length
* 61045 61568: contig of 524 bp in length
* 61569 61648: gap of unknown length
* 61649 62198: contig of 550 bp in length
* 62199 62278: gap of unknown length
* 62279 62868: contig of 590 bp in length
* 62869 62948: gap of unknown length
* 62949 63560: contig of 612 bp in length
* 63561 63640: gap of unknown length
* 63641 64213: contig of 573 bp in length
* 64214 64293: gap of unknown length
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* 65583 65662: gap of unknown length
* 65663 66213: contig of 551 bp in length

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  Quality: 40.00      Length: 9
  Ratio: 4.444       Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 88.889

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alignment_block:

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US-08-653-294-11 x AC009355/rev ...
Align seg 1/1 to reverse of: AC009355 from: 1 to: 66958

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1 TyrArgLeuLeuLeuArgLeuAnGlu 9
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46014 TTCCGGTACTTATTTCGGTTGACGAG 45988

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seq_name: gb_pr3:AC005365

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seq_documentation_block:
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DEFINITION Homo sapiens chromosome 16, P1 clone 79-2A (LANL), complete
sequence.
ACCESSION AC005365
VERSION AC005365.1 GI:3367509
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homnidae; Homo.

```

```

REFERENCE
AUTHORS
Ricke D.O., Bruce D., Mundt M., Doggett N., Munk C., Saunders E.,
Robinson D., Jones M., Buckingham J., Chasteen L., Thompson S.,
Goodwin L., Bryant J., Tesmer J., Meincke L., Longmire J.,
White S., Ueng S., Tatum O., Campbell C., Fawcett J., Maltbie M.,
Misra M. and Deaven L.
Sequencing of Human Chromosome 16

```

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TITLE
JOURNAL
REFERENCE
2 (bases 1 to 86130)
AUTHORS
Ricke D.O.
Large Scale Sequence Analysis and Annotation with the Sequence
Comparison Analysis (SCAN) System

```

JOURNAL

```

3 (bases 1 to 86130)
REFERENCE
AUTHORS
Ricke D.O., Bruce D., Mundt M., Doggett N., Munk C., Saunders E.,
Robinson D., Jones M., Buckingham J., Chasteen L., Thompson S.,
Goodwin L., Bryant J., Tesmer J., Meincke L., Longmire J.,
White S., Ueng S., Tatum O., Campbell C., Fawcett J., Maltbie M.,
Misra M. and Deaven L.
Direct Submission
Submitted (01-AUG-1998) Center for Human Genome Studies, DOE Joint
Genome Institute, Los Alamos National Laboratory, MS M888, Los
Alamos, NM 87545, USA

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FEATURES

source

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Location/Qualifiers
1..86130

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repeat_region

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7853..8448
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misc_feature

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misc_feature

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repeat_region

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repeat_region

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repeat_region

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repeat_region

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repeat_region

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repeat_region

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repeat_region

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29408..29459
/rpt_family="MIR"

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repeat_region

```

30146..30276
/rpt_family="Alu"

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misc_feature

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30513..30631
note="GRAIL 2 excellent exon, frame 1"

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repeat_region

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30919..31186
/rpt_family="Alu"

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repeat_region

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31608..32152

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repeat_region /rpt_family="Alu" complement(32560..32851)
repeat_region /rpt_family="Alu" 33192..33293
misc_feature 35811..35929
repeat_region /note="GRAIL 2 excellent exon, frame 2"
36035..36312
repeat_region /rpt_family="Alu" complement(36626..36748)
repeat_region /rpt_family="L1" complement(37173..37468)
repeat_region 38519..38818
repeat_region /rpt_family="Alu" 40556..40673
repeat_region /rpt_family="MIR" 41506..41532
repeat_region /note="(A)27"
/rpt_type=tandem
/rpt_unit=A
repeat_region 45596..45892
repeat_region /rpt_family="Alu" 46881..46366
misc_feature 46477..46721
repeat_region /note="GRAIL 2 excellent exon, frame 0"
47484..47760
misc_feature 50106..50312
repeat_region /note="GRAIL 2 excellent exon, frame 2"
50512..50788
repeat_region /rpt_family="Alu" 50789..51068
repeat_region complement(52275..52538)
repeat_region /rpt_family="Alu" complement(53058..53720)
repeat_region 54695..54976
repeat_region /rpt_family="Alu" 55556..55829
repeat_region /rpt_family="Alu" 56365..56848
repeat_region /rpt_family="Alu" complement(57163..57314)
misc_feature /note="GRAIL 2 excellent exon, frame 1"
58926..59183
repeat_region /rpt_family="Alu" 59490..60014
repeat_region /rpt_family="Alu" complement(60362..60655)
repeat_region 60988..61273
repeat_region /rpt_family="Alu" 61417..61757
repeat_region /rpt_family="Alu" complement(62028..62141)
repeat_region complement(63160..63454)
repeat_region /rpt_family="Alu" complement(63604..63875)
repeat_region /rpt_family="Alu" complement(63894..64098)
misc_feature 64463..64588
repeat_region /note="GRAIL 2 excellent exon, frame 2"
64707..64992
misc_feature 66355..66492
repeat_region /note="GRAIL 2 excellent exon, frame 0"
complement(66679..66796)
/rpt_family="Alu"

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repeat_region 6995..67262
/rpt_family="Alu" 67741..68036
repeat_region /rpt_family="Alu" complement(69213..69456)
repeat_region /rpt_family="Alu" 70077..70906
repeat_region /rpt_family="L1" 71602..72045
repeat_region /rpt_family="L1" 72758..72876
repeat_region /rpt_family="MER46" complement(72760..72985)
repeat_region /rpt_family="MER46" complement(73181..73243)
repeat_region /rpt_family="MIR" complement(74246..74532)
repeat_region /rpt_family="Alu" 74601..74899
repeat_region /rpt_family="Alu" 74915..75035
repeat_region /rpt_family="L1" 75843..76122
repeat_region /rpt_family="Alu" 76105..76145
repeat_region /note="(A)41"
/rpt_type=tandem
/rpt_unit=A
repeat_region 76163..76652
repeat_region /rpt_family="Alu" 76635..76655
/rpt_family="Alu"

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alignment_scores:
  Quality: 40.00 Length: 10
  Ratio: 4.444 Gaps: 0
  Percent Similarity: 90.000 Percent Identity: 80.000

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alignment_block:

US-08-653-294-11 x AC005365 ..

Align seg 1/1 to: AC005365 from: 1 to: 86130

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10

|||||||:||||||| |||
23775 TACAGACTGTGCATCAGATTGAATCCCGAGA 23804

seq_name: gb_htg1:HS1141E20

seq_documentation_block:

LOCUS HS1141E20 97906 bp DNA HTG 23-NOV-1999
DEFINITION Homo sapiens chromosome 6 clone RP5-1141E20, *** SEQUENCING IN
PROGRESS ***, in unordered pieces.

ACCESSION AL109912

VERSION AL109912.4 GI:5870369

KEYWORDS HTG; HTGS_PHASE1.

SOURCE human.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 97906)

Direct Submission

Submitted (10-SEP-1999) Wellcome Trust Genome Campus, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquires:

humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk

On Sep 12, 1999 this sequence version replaced gi:5777438.

IMPORTANT: This sequence is unfinished and does not necessarily

represent the correct sequence. Work on the sequence is in progress

and the release of this data is based on the understanding that the

sequence may change as work continues. The sequence may be

contaminated with foreign sequence from E.coli, yeast, vector,

phage etc. Order of segments is not known; 800 n's separate

segments. Unfinished: dJ1141E20 Contig_ID: 00340 acc=AL109912
 Length: 64235 bp Unfinished: dJ1141E20 Contig_ID: 00618
 acc=AL109912 Length: 18869 bp Unfinished: dJ1141E20 Contig_ID:
 00679 acc=AL109912 Length: 13202 bp.

* NOTE: This is a 'working draft' sequence.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

FEATURES

Location/Qualifiers
 1. 97906
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="6"
 /clone="RP5-1141E20"
 /clone.lib="RPCI-5"
 BASE COUNT 30195 a 18298 c 18016 g 29795 t 1602 others
 ORIGIN

alignment_scores:

Quality: 40.00 Length: 10
 Ratio: 4.444 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-11 x HS1141E20/rev ..
 Align seg 1/1 to reverse of: HS1141E20 from: 1 to: 97906

1 TyrArgLeuIleArgLeuAsnGUArg 10

|||||||:::|||||||
 48886 TACAGACTTGTCATGATTAATCCAGA 48857

seq_name: gb_hgt3:AC008324

seq_documentation_block:
 LOCUS AC008324 122061 bp DNA HTG 06-AUG-1999
 DEFINITION Drosophila melanogaster chromosome 2 clone BACR25K01 (D854) RPCI-98
 25.K.1 map 25C-25D strain y; cn bw sp, *** SEQUENCING IN PROGRESS
 ***, 81 unordered pieces.

ACCESSION AC008324

VERSION AC008324.1 GI:5670415

KEYWORDS HTG; HTGS_PHASE1.

SOURCE fruit fly.

ORGANISM

Drosophila melanogaster
 Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
 Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE 1 (bases 1 to 122061)

AUTHORS
 Ceiniker,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
 Butenhoff,C., Champe,M., Chavez,C., Chev,M., Ciesiolka,L.,
 Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
 Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
 Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
 Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
 Pfeiffer,B., Poon,L., Sequelra,A., Sethi,H., Shrir,E.,
 Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
 Rubin,G.M.

Sequencing of Drosophila melanogaster

TITLE

JOURNAL

REFERENCE 2 (bases 1 to 122061)

AUTHORS
 Ceiniker,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
 Butenhoff,C., Champe,M., Chavez,C., Chev,M., Ciesiolka,L.,
 Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
 Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
 Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
 Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
 Pfeiffer,B., Poon,L., Sequelra,A., Sethi,H., Shrir,E.,
 Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
 Rubin,G.M.

Direct Submission

TITLE

JOURNAL

Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley

Laboratory, MS 84-121, Berkeley, CA 94720, USA

COMMENT

For further information about this sequence, including its location
 and relationship to other sequences, please visit our sequence
 archive web site (<http://www.fruitfly.org/sequence/>) or send email
 to bdg@fruitfly.berkeley.edu. All contigs in this submission meet
 the following cutoffs: length >= 200 bases.
 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 81 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1
 594: contig of 594 bp in length
 674: gap of unknown length
 1814: contig of 1140 bp in length
 1894: gap of unknown length
 2545: contig of 651 bp in length
 2625: gap of unknown length
 3091: contig of 466 bp in length
 3171: gap of unknown length
 4398: contig of 1227 bp in length
 4478: gap of unknown length
 5421: contig of 943 bp in length
 5501: gap of unknown length
 6196: contig of 695 bp in length
 6276: gap of unknown length
 7008: contig of 732 bp in length
 7088: gap of unknown length
 7770: contig of 882 bp in length
 7850: gap of unknown length
 8573: contig of 723 bp in length
 8653: gap of unknown length
 9411: contig of 758 bp in length
 9491: gap of unknown length
 10299: contig of 808 bp in length
 10379: gap of unknown length
 11647: contig of 1268 bp in length
 11727: gap of unknown length
 12578: contig of 851 bp in length
 12658: gap of unknown length
 13474: contig of 816 bp in length
 13554: gap of unknown length
 14454: contig of 900 bp in length
 14534: gap of unknown length
 15301: contig of 767 bp in length
 15381: gap of unknown length
 16820: contig of 1439 bp in length
 16900: gap of unknown length
 17817: contig of 917 bp in length
 17897: gap of unknown length
 19666: contig of 1769 bp in length
 19746: gap of unknown length
 21069: contig of 1323 bp in length
 21149: gap of unknown length
 22084: contig of 935 bp in length
 22164: gap of unknown length
 23539: contig of 1375 bp in length
 23619: gap of unknown length
 25170: contig of 1551 bp in length
 25250: gap of unknown length
 26158: contig of 908 bp in length
 26238: gap of unknown length
 27461: contig of 1223 bp in length
 27541: gap of unknown length
 28048: contig of 507 bp in length
 28128: gap of unknown length
 29477: contig of 1349 bp in length
 29557: gap of unknown length
 31107: contig of 1550 bp in length
 31187: gap of unknown length
 31856: contig of 669 bp in length
 31936: gap of unknown length

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* 31937 33142: contig of 1206 bp in length
* 33143 33222: gap of unknown length
* 33223 34713: contig of 1491 bp in length
* 34714 34793: gap of unknown length
* 34794 36055: contig of 1262 bp in length
* 36056 36135: gap of unknown length
* 36136 37830: contig of 1895 bp in length
* 37831 37910: gap of unknown length
* 37911 39370: contig of 1360 bp in length
* 39371 39350: gap of unknown length
* 39351 40199: contig of 849 bp in length
* 40200 40279: gap of unknown length
* 40280 42203: contig of 1924 bp in length
* 42204 42283: gap of unknown length
* 42284 44017: contig of 1734 bp in length
* 44018 44097: gap of unknown length
* 44098 45611: contig of 1514 bp in length
* 45612 45691: gap of unknown length
* 45692 47034: contig of 1343 bp in length
* 47035 47114: gap of unknown length
* 47115 48858: contig of 1744 bp in length
* 48859 48938: gap of unknown length
* 48939 50521: contig of 1583 bp in length
* 50522 50601: gap of unknown length
* 50602 52996: contig of 2395 bp in length
* 52997 53076: gap of unknown length
* 53077 54627: contig of 1551 bp in length
* 54628 54707: gap of unknown length
* 54708 56095: contig of 1388 bp in length
* 56096 56175: gap of unknown length
* 56176 57722: contig of 1547 bp in length
* 57723 57802: gap of unknown length
* 57803 60670: contig of 2868 bp in length
* 60671 60750: gap of unknown length
* 60751 62436: contig of 1686 bp in length
* 62437 62516: gap of unknown length
* 62517 64900: contig of 2384 bp in length
* 64901 64980: gap of unknown length
* 64981 67157: contig of 2177 bp in length
* 67158 67237: gap of unknown length
* 67238 70347: contig of 3110 bp in length
* 70348 70427: gap of unknown length
* 70428 73471: contig of 3044 bp in length
* 73472 73551: gap of unknown length
* 73552 76116: contig of 2565 bp in length
* 76117 76196: gap of unknown length
* 76197 79584: contig of 3388 bp in length
* 79585 79664: gap of unknown length
* 79665 82547: contig of 2883 bp in length
* 82548 82627: gap of unknown length
* 82628 85279: contig of 2652 bp in length
* 85280 85359: gap of unknown length
* 85360 88975: contig of 3516 bp in length
* 88976 88955: gap of unknown length
* 88956 93882: contig of 4927 bp in length
* 93883 93962: gap of unknown length
* 93963 98575: contig of 4613 bp in length
* 98576 98655: gap of unknown length
* 98656 105323: contig of 6668 bp in length
* 105324 105403: gap of unknown length
* 105404 106093: contig of 690 bp in length
* 106094 106173: gap of unknown length
* 106174 106901: contig of 728 bp in length
* 106902 106981: gap of unknown length
* 106982 107612: contig of 631 bp in length
* 107613 107692: gap of unknown length
* 107693 108426: contig of 734 bp in length
* 108427 108506: gap of unknown length
* 108507 109243: contig of 737 bp in length
* 109244 109323: gap of unknown length
* 109324 110081: contig of 758 bp in length
* 110082 110161: gap of unknown length
* 110162 110890: contig of 729 bp in length
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* 110891 110970: gap of unknown length
* 110971 111686: contig of 716 bp in length
* 111687 111766: gap of unknown length
* 111767 112474: contig of 708 bp in length
* 112475 112554: gap of unknown length
* 112555 113312: contig of 758 bp in length
* 113313 113392: gap of unknown length
* 113393 114092: contig of 700 bp in length
* 114093 114172: gap of unknown length
* 114173 114930: contig of 758 bp in length
* 114931 115010: gap of unknown length
* 115011 115817: contig of 807 bp in length
* 115818 115897: gap of unknown length
* 115898 115996: contig of 699 bp in length
* 115997 116766: gap of unknown length
* 116767 117456: contig of 780 bp in length
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alignment_scores:
  Quality: 40.00      Length: 9
  Ratio: 4.44         Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 88.889

alignment_block:
  US-08-653-294-11 x AC008324/rev ..

  Align seg 1/1 to reverse of: AC008324 from: 1 to: 122061
    1 TyArgLeuLeuIleArgLeuAsnGlu 9
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23076 TTCGGTACTTATCGGTGACGAG 23050
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seq_name: gb_pr3:HSJ1112D6

seq_documentation_block:

LOCUS HSJ1112D6 135305 bp DNA PRI 23-NOV-1999
DEFINITION Human DNA sequence from clone 1112D6 on chromosome 6q21-22.2,
complete sequence.

ACCESSION AL080317

VERSION AL080317.11 GI:5830430

KEYWORDS HTG; CpG Island.

SOURCE human

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 135305)

AUTHORS Patel,R.

TITLE Direct Submission

JOURNAL Submitted (09-SEP-1999) Sanger Centre, Hinxton, Cambridgeshire,

CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

requests: clonerequest@sanger.ac.uk

On Sep 6, 1999 this sequence version replaced gi:5791529.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

The following abbreviations are used to associate primary accession

numbers given in the feature table with their source databases:

Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WormPEP; Information

on the WormPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence is

the entire insert of clone 1112D6. This sequence has been finished

according to sequence map criteria as follows. An attempt is made

to resolve all sequencing problems, such as compressions and

repeats, but not necessarily within known annotated human repeat

sequence elements (e.g. Alu). Where the sequence is ambiguous,

there is an annotation using the 'unsure' feature key.

This sequence was generated from part of bacterial clone contigs of

human chromosome 6, constructed by the Sanger Centre Chromosome 6

Mapping Group. Further information can be found at

<http://www.sanger.ac.uk/HGP/Chr6>

1112b6 is from the library RPCI-5 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/VECTOR: pcypac2>.

FEATURES

source
1. 135305
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="6"
/map="q21-22.2"
/clone_lib="RPCI-5"
/clone="RP5-1112D6"
BASE COUNT 41171 a 26002 c 26091 g 42041 t
ORIGIN

alignment_scores:
Quality: 40.00 Length: 10
Ratio: 4.444 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-11 x HSJ1112D6/rev ..

Align seg 1/1 to reverse of: HSJ1112D6 from: 1 to: 135305

1 TyrArgLeuLeuileArgLeuAsnGluArg 10

|||||:|||||:|||||:|||||:|||||

99309 TACAGACTGTGCATCAGATTGAATCCCGA 99280

seq_name: gb_htg3:AC008327

seq_documentation_block:

LOCUS AC008327 185469 bp DNA HTG 20-AUG-1999
DEFINITION Drosophila melanogaster chromosome 2 clone BACR14K04 (D859), RPCI-98
14.K.4 map 27C-27C strain y; cn bw sp, *** SEQUENCING IN PROGRESS
*** 188 unordered pieces.

ACCESSION

AC008327

VERSION AC008327.2 GI:5748862

KEYWORDS HTG; HTGS_PHASE1.

SOURCE fruit fly.

ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

AUTHORS
Celniker,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
Butenhoff,C., Chape,M., Chavez,C., Chew,M., Ciesiolka,L.,
Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
Pfeiffer,B., Poon,L., Sequeira,A., Sethi,H., Snir,E.,
Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
Rubin,G.M.
Sequencing of Drosophila melanogaster
Unpublished
2 (bases 1 to 185469)
Celniker,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
Butenhoff,C., Chape,M., Chavez,C., Chew,M., Ciesiolka,L.,
Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
Pfeiffer,B., Poon,L., Sequeira,A., Sethi,H., Snir,E.,
Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
Rubin,G.M.

TITLE

Sequencing of Drosophila melanogaster

REFERENCE

AUTHORS
Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On Aug 20, 1999 this sequence version replaced gi:5670411.
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive web site (<http://www.fruitfly.org/sequence/>) or send email

TITLE

JOURNAL
Submitted (02-AUG-1999) Drosophila Genome Center, Lawrence Berkeley
Laboratory, MS 64-121, Berkeley, CA 94720, USA
On Aug 20, 1999 this sequence version replaced gi:5670411.
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive web site (<http://www.fruitfly.org/sequence/>) or send email

to bdg@fruitfly.berkeley.edu. All contigs in this submission meet
the following cutoffs: length >= 200 bases.

* NOTE: This is a 'working draft' sequence. It currently
* consists of 188 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 672: contig of 672 bp in length
* 673 752: gap of unknown length
* 753 1942: contig of 1190 bp in length
* 1943 2022: gap of unknown length
* 2023 2902: contig of 880 bp in length
* 2903 2982: gap of unknown length
* 2983 4320: contig of 1338 bp in length
* 4321 4400: gap of unknown length
* 4401 5184: contig of 784 bp in length
* 5185 5264: gap of unknown length
* 5265 6058: contig of 794 bp in length
* 6059 6138: gap of unknown length
* 6139 6857: contig of 719 bp in length
* 6858 6937: gap of unknown length
* 6938 8116: contig of 1179 bp in length
* 8117 8893: contig of 697 bp in length
* 8894 8973: gap of unknown length
* 8974 9623: contig of 650 bp in length
* 9624 9703: gap of unknown length
* 9704 10743: contig of 1041 bp in length
* 10745 10824: gap of unknown length
* 10825 11514: contig of 690 bp in length
* 11515 11594: gap of unknown length
* 11595 12380: contig of 786 bp in length
* 12381 12460: gap of unknown length
* 12461 13282: contig of 822 bp in length
* 13283 13282: gap of unknown length
* 13283 13363: contig of 576 bp in length
* 13363 14018: gap of unknown length
* 14019 15240: contig of 1222 bp in length
* 15241 15320: gap of unknown length
* 15321 16141: contig of 821 bp in length
* 16142 16221: gap of unknown length
* 16222 16940: contig of 719 bp in length
* 16941 17020: gap of unknown length
* 17021 17930: contig of 910 bp in length
* 17931 18010: gap of unknown length
* 18011 18792: contig of 782 bp in length
* 18793 18872: gap of unknown length
* 18873 19592: contig of 720 bp in length
* 19593 19672: gap of unknown length
* 19673 20826: contig of 1154 bp in length
* 20827 20906: gap of unknown length
* 20907 21710: contig of 804 bp in length
* 21711 21790: gap of unknown length
* 21791 22994: contig of 1204 bp in length
* 22995 23074: gap of unknown length
* 23075 24155: contig of 1081 bp in length
* 24156 24335: gap of unknown length
* 24336 25381: contig of 1146 bp in length
* 25382 25461: gap of unknown length
* 25462 26295: contig of 834 bp in length
* 26296 26375: gap of unknown length
* 26376 27866: contig of 1411 bp in length
* 27867 27866: gap of unknown length
* 27867 28593: contig of 827 bp in length
* 28594 28773: gap of unknown length
* 28774 29479: contig of 706 bp in length
* 29480 29559: gap of unknown length
* 29560 30139: contig of 580 bp in length
* 30140 30219: gap of unknown length
* 30220 30861: contig of 642 bp in length

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* 30862 30941: gap of unknown length
* 30942 32336: contig of 1395 bp in length
* 32337 32416: gap of unknown length
* 32417 32854: contig of 438 bp in length
* 32855 32934: gap of unknown length
* 32935 34121: contig of 1187 bp in length
* 34122 34201: gap of unknown length
* 34202 34933: contig of 732 bp in length
* 34934 35013: gap of unknown length
* 35014 35817: contig of 804 bp in length
* 35818 35897: gap of unknown length
* 35898 36647: contig of 750 bp in length
* 36648 36727: gap of unknown length
* 36728 37810: contig of 1083 bp in length
* 37811 37890: gap of unknown length
* 37891 39903: contig of 2013 bp in length
* 39904 39984: gap of unknown length
* 39984 41099: contig of 1116 bp in length
* 41100 41179: gap of unknown length
* 41180 42669: contig of 1490 bp in length
* 42670 42749: gap of unknown length
* 42750 44136: contig of 1387 bp in length
* 44137 44216: gap of unknown length
* 44217 45419: contig of 1203 bp in length
* 45420 45499: gap of unknown length
* 45500 46236: contig of 737 bp in length
* 46237 46316: gap of unknown length
* 46317 47980: contig of 1664 bp in length
* 47981 48060: gap of unknown length
* 48061 49473: contig of 1413 bp in length
* 49474 49553: gap of unknown length
* 49554 50637: contig of 1084 bp in length
* 50638 50717: gap of unknown length
* 50718 52416: contig of 1699 bp in length
* 52417 52496: gap of unknown length
* 52497 53367: contig of 871 bp in length
* 53368 53447: gap of unknown length
* 53448 54708: contig of 1261 bp in length
* 54709 54788: gap of unknown length
* 54789 56337: contig of 1549 bp in length
* 56338 56417: gap of unknown length
* 56418 57892: contig of 1475 bp in length
* 57893 57972: gap of unknown length
* 57973 59071: contig of 1099 bp in length
* 59072 59151: gap of unknown length
* 59152 60999: contig of 1848 bp in length
* 61000 61079: gap of unknown length
* 61080 62756: contig of 1677 bp in length
* 62757 62836: gap of unknown length
* 62837 64047: contig of 1211 bp in length
* 64048 64127: gap of unknown length
* 64128 65337: contig of 1210 bp in length
* 65338 65417: gap of unknown length
* 65418 67030: contig of 1613 bp in length
* 67031 67110: gap of unknown length
* 67111 68452: contig of 1342 bp in length
* 68453 68532: gap of unknown length
* 68533 70469: contig of 1937 bp in length
* 70470 70549: gap of unknown length
* 70550 71670: contig of 1121 bp in length
* 71671 71750: gap of unknown length
* 71751 73015: contig of 1265 bp in length
* 73016 73095: gap of unknown length
* 73096 74102: contig of 1007 bp in length
* 74103 74182: gap of unknown length
* 74183 75541: contig of 1359 bp in length
* 75542 75621: gap of unknown length
* 75622 77100: contig of 1479 bp in length
* 77101 77181: gap of unknown length
* 77182 79860: contig of 2680 bp in length
* 79861 79940: gap of unknown length
* 79941 80707: contig of 767 bp in length
* 80708 80787: gap of unknown length
```

```
*
* 80788 83328: contig of 2541 bp in length
* 83329 83408: gap of unknown length
* 83409 85899: contig of 2491 bp in length
* 85900 85979: gap of unknown length
* 85980 88101: contig of 2122 bp in length
* 88102 88181: gap of unknown length
* 88182 90444: contig of 2163 bp in length
* 90445 90424: gap of unknown length
* 90425 94408: contig of 3984 bp in length
* 94409 94488: gap of unknown length
* 94489 99242: contig of 4754 bp in length
* 99243 99322: gap of unknown length
* 99323 99919: contig of 597 bp in length

alignment_scores:
  Quality: 40.00      Length: 9
  Ratio: 4.444       Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 88.889

alignment_block:
  US-08-653-294-11 x AC008327/rev ..
  Align seg 1/1 to reverse of: AC008327 from: 1 to: 185469

  1 TyrArgLeuLeuIleArgLeuAsnGlu 9
  ::::::::::::::::::::::::::::::
  161633 TTCCGGTTACTTATTCGTTGAACGAG 161607

seq_name: gb_htg7:AC017912

seq_documentation_block:
  LOCUS AC017912 49261 bp DNA HTG 09-DEC-1999
  DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered
  pieces.
  ACCESSION AC017912
  VERSION AC017912.1 GI:6553278
  KEYWORDS Htg; Hrgs_PHASE2.
  SOURCE fruit fly.
  ORGANISM Drosophila melanogaster
    Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
    Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
    Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
  REFERENCE 1 (bases 1 to 49261)
  AUTHORS Adams,M. and Venter,J.C.
  TITLE Direct Submission
  JOURNAL Submitted (09-DEC-1999) Celera Genomics, 45 West Gude Drive,
  Rockville, MD, USA
  COMMENT This sequence was identified as CDM:10212607 by the submitter.
  For more information on this record e-mail to fly@celera.com.
  * NOTE: This is a 'working draft' sequence.
  * This sequence will be replaced
  * by the finished sequence as soon as it is available and
  * the accession number will be preserved.

FEATURES
  source
  1..49261
  /organism="Drosophila melanogaster"
  /db_xref="taxon:7227"
  BASE COUNT 14872 a 9975 c 9922 g 14492 t
  ORIGIN

alignment_scores:
  Quality: 39.00      Length: 10
  Ratio: 3.900       Gaps: 0
  Percent Similarity: 100.000  Percent Identity: 70.000

alignment_block:
  US-08-653-294-11 x AC017912/rev ..
  Align seg 1/1 to reverse of: AC017912 from: 1 to: 49261

  1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
  ::::::::::::::::::::::::::::::
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/rpt_family="ALU"
complement(32617. .32681)
/rpt_family="ALU"
35367. .35529
/genes="WNT2"
/notes="match to human EST T29432 (NID:611530)"
complement(3854. .39080)
/rpt_family="ALU"
43070. .45031
/notes="CpG island (%GC=64.4, o/e=0.70, #CpGs=157)"
47242. .47360
/rpt_family="ALU"
complement(48223. .48514)
/rpt_family="ALU"
48717. .48995
/rpt_family="ALU"
complement(49005. .49032)
/rpt_family="L1"
51331. .51772
/rpt_family="L1"
52336. .52494
/rpt_family="L1"
52528. .52987
/rpt_family="L1"
54271. .54366
/rpt_family="L1"
55051. .55074
/rpt_family="L1"
55654. .55791
/rpt_family="L1"
55902. .55935
/rpt_family="L1"
56087. .56189
/rpt_family="L1"
complement(56256. .56548)
/rpt_family="ALU"
56855. .56695
/rpt_family="L1"
56730. .56802
/rpt_family="L1"
56818. .56938
/rpt_family="ALU"
57144. .57227
/rpt_family="L1"
complement(59426. .59717)
/rpt_family="ALU"
complement(60815. .61106)
/rpt_family="ALU"
complement(61569. .61869)
/rpt_family="ALU"
complement(63011. .67613)
/rpt_family="L1"
63980. .64398
/rpt_family="L1"
67964. .68015
/rpt_family="MER"
68059. .69083
/rpt_family="MER"
72312. .72350

alignment_scores:
  Quality: 39.00 Length: 10
  Ratio: 4.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x AC002465
Align seg 1/1 to: AC002465 from: 1 to: 155881
1 TyrArgLeuLeuAargLeuAsnGluArg 10
||||| : : : : :
15573 TATGACTTCTGTGAGTCAATGAGAGA 15602
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seq_name: gb_htg5:AC015280
seq_documentation_block:
LOCUS AC015280 159468 bp DNA HTG 16-NOV-1999
DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered
pieces.
ACCESSION AC015280
VERSION AC015280.1 GI:6436055
KEYWORDS HTG; HTGS_PHASE2.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 159468)
AUTHORS Adams, M. and Venter, J.C.
TITLE Direct Submission
JOURNAL Submitted (16-NOV-1999) Celera Genomics, 45 West Gude Drive,
Rockville, MD, USA
COMMENT This sequence was identified as CDM:10213506 by the submitter.
For further information on this sequence e-mail to fly@celera.com.
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
FEATURES
  source
    1. .159468
    /organism="Drosophila melanogaster"
    /db_xref="taxon:7227"
BASE COUNT 47335 a 32541 c 32291 g 47301 t
ORIGIN

alignment_scores:
  Quality: 39.00 Length: 10
  Ratio: 3.900 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x AC015280
Align seg 1/1 to: AC015280 from: 1 to: 159468
1 TyrArgLeuLeuAargLeuAsnGluArg 10
||||| : : : : :
44309 TATAAATTATTGATCAGGATCAATAGCCGA 44338

seq_name: gb_htg4:AC007054
seq_documentation_block:
LOCUS AC007054 171979 bp DNA HTG 13-OCT-1999
DEFINITION Drosophila melanogaster chromosome 2 clone BACR45018 (P527) RPCI-98
45.0.18 map 41E-41E strain y; cn bw sp, *** SEQUENCING IN PROGRESS
***, 13 unordered pieces.
ACCESSION AC007054
VERSION AC007054.22 GI:6041715
KEYWORDS HTG; HTGS_PHASE1.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 171979)
AUTHORS Celnikier, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazej, R.G.,
Butenhoif, C., Champagne, M., Chavez, C., Chew, M., Ciesiolka, L.,
Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L.,
Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L.,
Kim, E., Lee, B., Lewis, S., Li, P., Lomotan, M.A., Mazda, P.,
Moshrefi, A.R., Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S.,
Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E.,
Svirskas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and
Rubin, G.M.
```

TITLE
JOURNAL
REFERENCE
AUTHORS

Sequencing of Drosophila melanogaster

Unpublished

2 (bases 1 to 171979)

Celniker, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazej, R.G., Butenhoff, C., Champe, M., Chavez, C., Chew, M., Ciesiolka, L., Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L., Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L., Kim, E., Lee, B., Lewis, S., Li, P., Lomotan, M.A., Mazda, P., Moshrefi, A.R., Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S., Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E., Svirkas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and Rubin, G.M.

Direct Submission

Submitted (10-MAR-1999) Drosophila Genome Center, Lawrence Berkeley Laboratory, MS 64-121, Berkeley, CA 94720, USA

On Oct 15, 1999 this sequence version replaced gi:5922045.

For further information about this sequence, including its location and relationship to other sequences, please visit our sequence archive Web site (<http://www.fruitfly.org/sequence/>) or send email to bdg@fruitfly.berkeley.edu. All contigs in this submission meet the following cutoffs: length >= 200 bases.

* NOTE: This is a 'working draft' sequence. It currently consists of 13 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 551: contig of 651 bp in length
652 731: gap of unknown length
732 1409: contig of 678 bp in length
1410 1489: gap of unknown length
1490 1840: contig of 351 bp in length
1841 1920: gap of unknown length
1921 2426: contig of 506 bp in length
2427 2506: gap of unknown length
2507 71748: contig of 69242 bp in length
71749 167344: contig of 95416 bp in length
167345 167324: gap of unknown length
167325 167942: contig of 618 bp in length
167943 168022: gap of unknown length
168023 168812: contig of 790 bp in length
168813 168892: gap of unknown length
168893 169544: contig of 652 bp in length
169545 169524: gap of unknown length
169525 169896: contig of 272 bp in length
169897 169876: gap of unknown length
169877 170314: contig of 338 bp in length
170315 170394: gap of unknown length
170395 171415: contig of 1021 bp in length
171416 171495: gap of unknown length
171496 171979: contig of 484 bp in length.

Location/Qualifiers

1..171979
/organism="Drosophila melanogaster"
/strain="Y: cn bw sp"
/db_xref="taxon:7227"
/chromosome="2"
/map="41E-41E"
/clone="BACR45018 (D527) RPCI-98 45.0.18"
/clone_lib="RPCI-98 (Roswell Park Cancer Institute Drosophila melanogaster BAC library, partial EcoRI in PBAC3.6")

BASE COUNT 50278 a 35745 c 35444 g 49502 t 1010 others

ORIGIN

alignment_scores:
Quality: 39.00 Length: 10
Ratio: 3.900 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-11 x AC007054/rev ..

Align seg 1/1 to reverse of: AC007054 from: 1 to: 171979

1 TTTArGLeuLeuLeuArgLeuAnGluArg 10
||||:|||||:|||||:|||||:|||||:|||||
128523 TATAAACTTGTGATCAGGATCAATAGCCGA 128494

seq_name: gb_htg4:AC006467

seq_documentation_block:

LOCUS AC006467 175695 bp DNA HTG 27-OCT-1999
DEFINITION Drosophila melanogaster chromosome 2 clone BACR03108 (D532) RPCI-98
03.L.8 map 40A-40C strain Y; cn bw sp, *** SEQUENCING IN PROGRESS
*** 9 unordered pieces.

ACCESSION AC006467.11 GI:6136329

VERSION HTG; HTGS_PHASE1.

KEYWORDS fruit fly.

ORGANISM

Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

1 (bases 1 to 175695)

AUTHORS

Celniker, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazej, R.G., Butenhoff, C., Champe, M., Chavez, C., Chew, M., Ciesiolka, L., Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L., Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L., Kim, E., Lee, B., Lewis, S., Li, P., Lomotan, M.A., Mazda, P., Moshrefi, A.R., Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S., Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E., Svirkas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and Rubin, G.M.

TITLE
JOURNAL
REFERENCE

Unpublished

2 (bases 1 to 175695)

AUTHORS

Celniker, S.E., Agbayani, A., Arcaina, T.T., Baxter, E., Blazej, R.G., Butenhoff, C., Champe, M., Chavez, C., Chew, M., Ciesiolka, L., Doyle, C.M., Farfan, D.E., Galle, R., George, R.A., Harris, N.L., Hoskins, R.A., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L., Kim, E., Lee, B., Lewis, S., Li, P., Lomotan, M.A., Mazda, P., Moshrefi, A.R., Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S., Pfeiffer, B., Poon, L., Sequeira, A., Sethi, H., Snir, E., Svirkas, R.R., Wan, K.H., Weinburg, T., Zhang, R., Zieran, L.L. and Rubin, G.M.

Direct Submission

Submitted (29-JAN-1999) Drosophila Genome Center, Lawrence Berkeley Laboratory, MS 64-121, Berkeley, CA 94720, USA

On Oct 27, 1999 this sequence version replaced gi:5670661.
For further information about this sequence, including its location and relationship to other sequences, please visit our sequence archive Web site (<http://www.fruitfly.org/sequence/>) or send email to bdg@fruitfly.berkeley.edu. All contigs in this submission meet the following cutoffs: length >= 200 bases.

* NOTE: This is a 'working draft' sequence. It currently consists of 9 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 550: contig of 550 bp in length
551 630: gap of unknown length
631 1229: contig of 599 bp in length
1230 1309: gap of unknown length
1310 1869: contig of 560 bp in length
1870 1949: gap of unknown length
1950 3022: contig of 1073 bp in length
3023 3102: gap of unknown length
3103 7823: contig of 4721 bp in length


```

* 7824 7903: gap of unknown length
* 7904 173453: contig of 16550 bp in length
* 173454 173533: gap of unknown length
* 173534 174264: contig of 731 bp in length
* 174265 174344: gap of unknown length
* 174345 174802: contig of 458 bp in length
* 174803 174882: gap of unknown length
* 174883 175695: contig of 813 bp in length.
FEATURES
    Location/Qualifiers
    1..175695
    /organism="Drosophila melanogaster"
    /strain="y; cn bw sp"
    /db_xref="taxon:7227"
    /chromosome="2"
    /map="40A-40C"
    /clone="BACR03L08 (D532) RPCI-98 03.L.8"
    /clone_lib="RPCI-98 (Roswell Park Cancer Institute
Drosophila melanogaster BAC library, partial EcoRI in
PRACE3.6"
BASE COUNT 51759 a 35773 c 35603 g 51919 t 641 others
ORIGIN

alignment_scores:
    Quality: 39.00      Length: 10
    Ratio: 3.900      Gaps: 0
    Percent Similarity: 100.000      Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x AC006467 ..

Align seg 1/1 to: AC006467 from: 1 to: 175695

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10
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48184 TATAAATTATGTCAGGATCAATAGCCGA 48213

seq_name: gb_hgt3:AC008682

seq_documentation_block:
LOCUS AC008682 216649 bp DNA HTG 03-AUG-1999
DEFINITION Homo sapiens chromosome 5 clone CIT978SKB_54G2, *** SEQUENCING IN
PROGRESS ***, 63 unordered pieces.
ACCESSION AC008682
VERSION AC008682.1 GI:5685896
KEYWORDS HTG; HTGS_PHASE1.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
  1 (bases 1 to 216649)
  AUTHORS DOE Joint Genome Institute.
  TITLE Sequencing of Human Chromosome 5
  JOURNAL Unpublished
REFERENCE
  2 (bases 1 to 216649)
  DOE Joint Genome Institute.
  Direct Submission
  TITLE Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
  JOURNAL Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
  www.jgi.doe.gov.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 63 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence.
* as soon as it is available and the accession number will
* be preserved.
* 1 637: contig of 637 bp in length
* gap of unknown length
* 538 1631: contig of 994 bp in length
* gap of unknown length
* 1632 2553: contig of 922 bp in length

```

```

* 2554 3319: gap of unknown length
* contig of 766 bp in length
* 3320 3994: contig of 675 bp in length
* gap of unknown length
* 3995 4648: contig of 654 bp in length
* gap of unknown length
* 4649 5512: contig of 864 bp in length
* gap of unknown length
* 5513 6208: contig of 696 bp in length
* gap of unknown length
* 6209 6922: contig of 714 bp in length
* gap of unknown length
* 6923 8025: contig of 1103 bp in length
* gap of unknown length
* 8026 9274: contig of 1249 bp in length
* gap of unknown length
* 9275 10498: contig of 1224 bp in length
* gap of unknown length
* 10499 11634: contig of 1136 bp in length
* gap of unknown length
* 11635 12332: contig of 698 bp in length
* gap of unknown length
* 12333 13945: contig of 1613 bp in length
* gap of unknown length
* 13946 15278: contig of 1333 bp in length
* gap of unknown length
* 15279 16574: contig of 1296 bp in length
* gap of unknown length
* 16575 17988: contig of 1414 bp in length
* gap of unknown length
* 17989 19075: contig of 1087 bp in length
* gap of unknown length
* 19076 20518: contig of 1443 bp in length
* gap of unknown length
* 20519 21769: contig of 1251 bp in length
* gap of unknown length
* 21770 22593: contig of 824 bp in length
* gap of unknown length
* 22594 23584: contig of 991 bp in length
* gap of unknown length
* 23585 24947: contig of 1363 bp in length
* gap of unknown length
* 24948 26399: contig of 1452 bp in length
* gap of unknown length
* 26400 27550: contig of 1151 bp in length
* gap of unknown length
* 27551 28267: contig of 717 bp in length
* gap of unknown length
* 28268 29109: contig of 842 bp in length
* gap of unknown length
* 29110 30110: contig of 1001 bp in length
* gap of unknown length
* 30111 31414: contig of 1304 bp in length
* gap of unknown length
* 31415 32809: contig of 1395 bp in length
* gap of unknown length
* 32810 34507: contig of 1698 bp in length
* gap of unknown length
* 34508 36466: contig of 1959 bp in length
* gap of unknown length
* 36467 38262: contig of 1796 bp in length
* gap of unknown length
* 38263 39740: contig of 1478 bp in length
* gap of unknown length
* 39741 40973: contig of 1233 bp in length
* gap of unknown length
* 40974 43125: contig of 2152 bp in length
* gap of unknown length
* 43126 46386: contig of 3261 bp in length
* gap of unknown length
* 46387 48948: contig of 2562 bp in length
* gap of unknown length

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* 48949 50932: contig of 1984 bp in length
* 50933 gap of unknown length
* 54161: contig of 3229 bp in length
* 54162 gap of unknown length
* 54164: contig of 2253 bp in length
* 54165 gap of unknown length
* 54166: contig of 4198 bp in length
* 54167 gap of unknown length
* 54168: contig of 1797 bp in length
* 54169 gap of unknown length
* 54170: contig of 4476 bp in length
* 54171 gap of unknown length
* 54172: contig of 3238 bp in length
* 54173 gap of unknown length
* 54174: contig of 4411 bp in length
* 54175 gap of unknown length
* 54176: contig of 3681 bp in length
* 54177 gap of unknown length
* 54178: contig of 4087 bp in length
* 54179 gap of unknown length
* 54180: contig of 3311 bp in length
* 54181 gap of unknown length
* 54182: contig of 2161 bp in length
* 54183 gap of unknown length
* 54184: contig of 3911 bp in length
* 54185 gap of unknown length
* 54186: contig of 7702 bp in length
* 54187 gap of unknown length
* 54188: contig of 4273 bp in length
* 54189 gap of unknown length
* 54190: contig of 6290 bp in length
* 54191 gap of unknown length
* 54192: contig of 9840 bp in length
* 54193 gap of unknown length
* 54194: contig of 6733 bp in length
* 54195 gap of unknown length
* 54196: contig of 5007 bp in length
* 54197 gap of unknown length
* 54198: contig of 7400 bp in length
* 54199 gap of unknown length
* 54200: contig of 10850 bp in length
* 54201 gap of unknown length
* 54202: contig of 15401 bp in length
* 54203 gap of unknown length
* 54204: contig of 16904 bp in length
* 54205 gap of unknown length
* 54206: contig of 34564 bp in length
* 54207 Location/Qualifiers
* 54208 1..216649
* 54209 /organism="Homo sapiens"
* 54210 /db_xref="taxon:9606"
* 54211 /chromosome="5"
* 54212 /clone="CIT9785KB_54G2"
* 54213 BASE COUNT 66797 a 42466 c 41332 g 66015 t 39 others
* 54214 ORIGIN
```

FEATURES

source

```
alignment_scores:
  Quality: 39.00 Length: 10
  Ratio: 3.900 Gaps: 0
  Percent Similarity: 100.000 Percent Identity: 70.000
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alignment_block:

US-08-653-294-11 x AC008682/rev ..

Align seg 1/1 to reverse of: AC008682 from: 1 to: 216649

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10

|||||:|||||:|||||:|||||:|||||

5329 TATAGGCTATTATAAGGATAATAATCGG 5300

OM of: US-08-653-294-11 to: N_Geneseq_36.* out_format : pfs

Date: Feb 8, 2000 1:27 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

-MODEL=framet_p2n.model -DEV=xlp
-O=/cgn1_1/USPTO.spool/US08653294/runat_04022000_160701_15807/app_query.fasta.1
-DB=N_Geneseq_36 -OPMT=fastap -SUFFIX=ring -GAPOP=12.000
-GAPEXT=4.000 -MINMATCH=0.100 -LOPCL=0.000 -LOOPEXT=0.000
-GAPOP=4.500 -GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500
-XGAPOP=6.000 -XGAPEXT=7.000 -XGAPOP=10.000 -XGAPEXT=0.500
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=blomsum62
-TRANS=human40.cdi -LIST=45 -DOCLIGN=200 -THR_SCORE=pct
-ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -MINLEN=0
-MAXLEN=100000 -USER=US08653294 -NCPU=6 -ICPU=3 -NO_XLPXY -WAIT
-THREADS=1

Search information block:

Query: US-08-653-294-11

Query length: 10

Database: N_Geneseq_36.*

Database sequences: 311585

Database length: 125096042

Search time (sec): 590.520000

score_list:

Sequence	Strd Orig	ZScore	EScore	Len	Documentation
N_Geneseq_36:T29774	- 37.00	115.12	82.10	2040	Bacillus thuringiensis ssp. is
N_Geneseq_36:N90712	- 37.00	114.87	84.82	2100	! cryd gene. Bacillus thuringie
N_Geneseq_36:Q14809	- 37.00	112.34	117.32	2802	! Btm PG14 72kDa Cry insecticida
N_Geneseq_36:Q14810	- 37.00	112.04	121.99	2901	! Btm PG14 72kDa Cry insecticida
N_Geneseq_36:V84560	+ 35.00	108.51	191.66	1842	! Human secreted protein gene 15
N_Geneseq_36:X20591	- 35.00	102.03	440.16	3858	! Polynucleotide sequence from t
N_Geneseq_36:V65599	+ 35.00	99.34	621.49	5243	! Fowlpox virus vector DNA fragm
N_Geneseq_36:T20013	+ 34.00	120.47	41.34	307	! Human gene signature HUMG01151
N_Geneseq_36:T77268	+ 34.00	118.29	54.74	394	! Staphylococcus aureus contig SB
N_Geneseq_36:X18185	+ 34.00	111.97	123.10	810	! S. aureus gldB coding sequence.
N_Geneseq_36:X18186	+ 34.00	111.04	138.59	900	! S. aureus gldB ORF sequence. Ne
N_Geneseq_36:V82078	+ 34.00	107.82	209.57	1300	! DNA encoding a partial gldA H
N_Geneseq_36:T74973	+ 34.00	107.18	227.42	1398	! Staphylococcus aureus contig S
N_Geneseq_36:Q20995	+ 34.00	106.22	257.27	1560	! EH-4 GC gene. Nucleic acid se
N_Geneseq_36:T82077	+ 34.00	103.48	365.56	2132	! DNA encoding a gldA protein.
N_Geneseq_36:V07921	- 34.00	102.67	405.51	2338	! Helicobacter pylori 76 kDa pol
N_Geneseq_36:V07916	- 34.00	102.34	423.30	2429	! Helicobacter pylori 76 kDa pol
N_Geneseq_36:V74500	+ 34.00	96.30	917.45	4832	! Staphylococcus aureus contig S
N_Geneseq_36:X20560	+ 34.00	91.39	1.7e+03	8467	! Polynucleotide sequence from t
N_Geneseq_36:T74420	+ 34.00	84.85	4.0e+03	17846	! Staphylococcus aureus contig
N_Geneseq_36:V48249	+ 33.00	103.71	354.70	1353	! Nucleotide sequence encoding s
N_Geneseq_36:T74701	- 33.00	101.27	485.32	1788	! Staphylococcus aureus contig S
N_Geneseq_36:T74662	- 33.00	101.13	493.87	1816	! Staphylococcus aureus contig S
N_Geneseq_36:X13313	+ 33.00	98.90	658.07	2344	! Staphylococcus aureus contig S
N_Geneseq_36:T42751	+ 33.00	89.81	2.1e+03	6608	! Chicken CHD-1A gene. Avian chr
N_Geneseq_36:V20767	+ 33.00	86.26	3.3e+03	9898	! Human OCIF genome DNA-2. Inhib
N_Geneseq_36:T33183	+ 33.00	86.01	3.4e+03	10190	! Fragment of human OCIF genom
N_Geneseq_36:X13065	+ 33.00	75.77	1.3e+04	32768	! Enterococcus faecalis genome
N_Geneseq_36:X13095	+ 32.50	79.92	7.5e+03	16484	! Enterococcus faecalis genome
N_Geneseq_36:Q39810	+ 32.00	112.00	122.63	343	! Expressed Sequence Tag human ge
N_Geneseq_36:Q59222	+ 32.00	112.00	122.63	343	! Human brain Expressed Sequence
N_Geneseq_36:T20671	+ 32.00	109.73	163.93	444	! Human gene signature HUMG01887
N_Geneseq_36:Q99805	+ 32.00	104.24	331.74	831	! Thaumatin like gene PR-5mz. New
N_Geneseq_36:Q70084	- 32.00	103.79	351.11	874	! Bacillus thuringiensis transcri
N_Geneseq_36:T84095	+ 32.00	102.34	423.26	1032	! DNA encoding a Staphylococcus
N_Geneseq_36:T75169	+ 32.00	101.19	490.23	1176	! Staphylococcus aureus contig S
N_Geneseq_36:T33136	- 32.00	99.76	588.77	1384	! Broccoli ACC synthase genomic
N_Geneseq_36:Q13581	- 32.00	99.64	598.35	1404	! A. altocetigenes membrane-bound
N_Geneseq_36:Q20384	- 32.00	99.64	598.35	1404	! ADH complex structural gene (4
N_Geneseq_36:T59378	- 32.00	98.78	668.28	1549	! 3' untranslated region of the
N_Geneseq_36:X02018	- 32.00	98.78	668.28	1549	! D. melanogaster tipE+ 4kb clon

N_Geneseq_36:V73474 - 32.00 97.61 775.93 1769 ! Mouse G3BP cDNA. New ubiqul
N_Geneseq_36:Q77686 + 32.00 96.17 933.97 2086 ! Temp. sensitive autolysing
N_Geneseq_36:T31725 - 32.00 95.99 955.65 2129 ! Human GAP-SH3 domain bindin
N_Geneseq_36:N91438 - 32.00 92.80 1.4e+03 3063 ! DNA encoding glycine rich p
seq_name: N_Geneseq_36:T29774

seq_documentation_block:

ID T29774 standard; DNA; 2040 BP.

AC T29774;

DT 19-NOV-1996 (first entry)

DE Bacillus thuringiensis ssp. israelensis CryIVD protein DNA.

KW CryIVD; toxic protein; crystal toxin; expression construct;

KW transformed cyanobacteria; phycoerythrin beta; cpCB; promoter;

OS insecticide; dipteran larvae; mosquito; blackfly; ss.

OS Bacillus thuringiensis.

FT Key

FT Location/Qualifiers

FT cds

FT US518897-A.

PD 21-MAY-1996.

PD 04-MAY-1992; 877876.

PR 04-MAY-1992; US-877876.

PR 28-JAN-1994; US-188581.

PA (OYME-) UNIV MEMPHIS STATE.

PI Murphy RC, Stevens SE;

DR WPI; 96-259063/26.

DR P-PSDB; R97735.

PT New DNA construct for expressing cryIV D protein in cyanobacteria

PT under control of a phycoerythrin beta promoter, useful for control of

PT dipteran larvae in water

PS Example 1; Columns 9-14; 20pp; English.

CC The present sequence encodes the B. thuringiensis ssp. israelensis

CC CryIVD toxic protein, which was used in the prep. of a claimed DNA

CC construct for the expression of CryIVD in cyanobacteria, comprising

CC the present sequence under the control of phycoerythrin beta (epCB)

CC promoter. Cyanobacteria (which may be adapted for growth in fresh

CC or brackish water) transformed with the construct can be used

CC as insecticides for controlling dipteran larvae, esp. those of

CC Culex pipiens (mosquito) larvae surviving after 4 days with

CC cyanobacteria transformed with the claimed DNA construct as their

CC only food source was 51 %, compared to 94 % for those fed with

CC cyanobacteria transformed with an empty plasmid. In the

CC cyanobacteria, CryIVD is efficiently expressed under the control of

CC the strong cpCB promoter, even though the CryIVD gene contains 19

CC AUA which are generally poorly translated (if at all) in

CC cyanobacteria.

CC Sequence 2040 BP; 716 A; 315 C; 373 G; 636 T;

alignment_scores:

Quality: 37.00 Length: 9

Ratio: 4.111 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-08-653-294-11 x T29774/rev ..

Align seg 1/1 to reverse of: T29774 from: 1 to: 2040

1 TyrArgLeuLeuLeuArgLeuAunGlu 9

|||||

542 TTCGCTCTTTTAAATGTAATGAA 516

seq_name: N_Geneseq_36:N90712

seq_documentation_block:

ID N90712 standard; DNA; 2100 BP.

AC N90712;

DT 09-JAN-1990 (first entry)

DE cryd gene.

KW cryd protein; Bacillus thuringiensis; biopesticide.

OS Bacillus thuringiensis var. israelensis.
 PN WO8907605-A.
 PD 24-AUG-1989.
 PF 17-FEB-1989; U00663.
 PR 19-FEB-1988; US-158176.
 PA (ECOG) Ecogen Inc.
 PI Donovan WP;
 DR WPI: 89-263682/36.
 DR P-PSDB: P91462.
 PT Bacillus thuringiensis var. israelensis cry D toxin gene and proteins
 PT - used for producing insecticide compsns. active against Dipteran species.
 PS Claim 1; fig 2; 58pp; English.
 CC cryD gene is inserted into plasmid and used to transform a microorganism.
 CC The 67kD protein encoded by the gene has insecticidal activity against
 CC dipteran larvae.
 SQ Sequence 2100 BP; 746 A; 316 C; 378 G; 660 T;

alignment_scores:
 Quality: 37.00 Length: 9
 Ratio: 4.111 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-08-653-294-11 x N90712/rev ..

Align seg 1/1 to reverse of: N90712 from: 1 to: 2100

1 TyrArgLeuLeuIleArgLeuAsnGlu 9

542 TTCGGTCTTTTAATAAGTAAATGAA 516

seq_name: N_Geneseq_36:Q14809

seq_documentation_block:

ID Q14809 standard; DNA; 2802 BP.
 AC Q14809;
 DT 10-FEB-1992 (first entry)
 DE Btm PG14 72kDa Cry insecticidal protein/25kDa Cyt A fusion gene.
 KW chimeric; fusion protein; insecticide; Lepidoptera larvae;
 KW midgut targeting; bacterial endotoxin; ss.
 OS Bacillus thuringiensis subspecies morrisoni.
 PN WO9117254-A.
 PD 14-NOV-1991.
 PF 02-MAY-1991; U03008.
 PR 03-MAY-1990; US-518575.
 PA (REGC) UNIV OF CALIFORNIA.
 PI Sivasubramanian N, Federici A;
 DR WPI: 91-353775/48.
 PT Extending host range or toxicity of insecticidal proteins - using
 PT protein capable of binding to gut epithelium of insects
 PS Claim 34; Fig 21; 61pp; English.
 CC This fusion gene comprises sequences isolated from the PG14 strain
 CC of B.thuringiensis subsp. morrisoni. The cytA gene encodes a
 CC cytolytic protein which has high affinity for the lipid portion of
 CC cell membranes. After ingestion by insects, the 27 kDa Cyt A protein
 CC is cleaved by midgut proteases to a relatively resistant core of 25
 CC kDa. The Cyt A coding sequence is combined with the Cry toxin, also
 CC from B.thuringiensis.
 SQ Sequence 2802 BP; 984 A; 395 C; 544 G; 879 T;

alignment_scores:
 Quality: 37.00 Length: 9
 Ratio: 4.111 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-08-653-294-11 x Q14809/rev ..

Align seg 1/1 to reverse of: Q14809 from: 1 to: 2802

1 TyrArgLeuLeuIleArgLeuAsnGlu 9

542 TTCGGTCTTTTAATAAGTAAATGAA 516

seq_name: N_Geneseq_36:Q14810

seq_documentation_block:

ID Q14810 standard; DNA; 2901 BP.
 AC Q14810;
 DT 10-FEB-1992 (first entry)
 DE Btm PG14 72kDa Cry insecticidal protein/27kDa Cyt A fusion gene.
 KW chimeric; fusion protein; insecticide; Lepidoptera larvae;
 KW midgut targeting; Cry A; bacterial endotoxin; ss.
 OS Bacillus thuringiensis subspecies morrisoni.
 PN WO9117254-A.
 PD 14-NOV-1991.
 PF 02-MAY-1991; U03008.
 PR 03-MAY-1990; US-518575.
 PA (REGC) UNIV OF CALIFORNIA.
 PI Sivasubramanian N, Federici A;
 DR WPI: 91-353775/48.
 PT Extending host range or toxicity of insecticidal proteins - using
 PT protein capable of binding to gut epithelium of insects
 PS Claim 33; Fig 22; 61pp; English.
 CC This fusion gene comprises sequences isolated from the PG14 strain
 CC of B.thuringiensis subsp. morrisoni. The cytA gene encodes a
 CC cytolytic protein which has high affinity for the lipid portion of
 CC cell membranes. After ingestion by insects, the 27 kDa Cyt A protein
 CC is cleaved by midgut proteases to a relatively resistant core of 25
 CC kDa. The Cyt A coding sequence is combined with the Cry toxin, also
 CC from B.thuringiensis.
 SQ Sequence 2901 BP; 1023 A; 462 C; 503 G; 913 T;

alignment_scores:

Quality: 37.00 Length: 9
 Ratio: 4.111 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-08-653-294-11 x Q14810/rev ..

Align seg 1/1 to reverse of: Q14810 from: 1 to: 2901

1 TyrArgLeuLeuIleArgLeuAsnGlu 9

542 TTCGGTCTTTTAATAAGTAAATGAA 516

seq_name: N_Geneseq_36:V84560

seq_documentation_block:

ID V84560 standard; DNA; 1842 BP.
 AC V84560;
 DT 01-MAR-1999 (first entry)
 DE Human secreted protein gene 150 clone HMSK035.
 KW Human; secreted protein; fusion protein; gene therapy; protein therapy;
 KW diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukaemia;
 KW developmental abnormality; foetal deficiency; blood; allergy; renal; ds;
 KW immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;
 KW inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;
 KW cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;
 KW osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;
 KW endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.
 OS Homo sapiens.
 PN WO9854963-A2.
 PD 10-DEC-1998.
 PF 04-JUN-1998; U11422.
 PR 18-DEC-1997; US-070923.
 PR 06-JUN-1997; US-048877.
 PR 06-JUN-1997; US-048881.
 PR 06-JUN-1997; US-048884.
 PR 06-JUN-1997; US-048893.
 PR 06-JUN-1997; US-048896.
 PR 06-JUN-1997; US-048899.

PR 06-JUN-1997; US-048915.
PR 06-JUN-1997; US-048949.
PR 06-JUN-1997; US-048964.
PR 06-JUN-1997; US-048972.
PR 06-JUN-1997; US-049020.
PR 06-JUN-1997; US-049375.
PR 05-SEP-1997; US-057628.
PR 05-SEP-1997; US-057635.
PR 05-SEP-1997; US-057644.
PR 05-SEP-1997; US-057647.
PR 05-SEP-1997; US-057650.
PR 05-SEP-1997; US-057661.
PR 05-SEP-1997; US-057667.
PR 05-SEP-1997; US-057761.
PR 05-SEP-1997; US-057764.
PR 05-SEP-1997; US-057770.
PR 05-SEP-1997; US-057775.
PR 05-SEP-1997; US-057778.
PR 06-JUN-1997; US-048875.
PR 06-JUN-1997; US-048878.
PR 06-JUN-1997; US-048882.
PR 06-JUN-1997; US-048885.
PR 06-JUN-1997; US-048894.
PR 06-JUN-1997; US-048897.
PR 06-JUN-1997; US-048900.
PR 06-JUN-1997; US-048916.
PR 06-JUN-1997; US-048962.
PR 06-JUN-1997; US-048970.
PR 06-JUN-1997; US-048974.
PR 06-JUN-1997; US-049373.
PR 05-SEP-1997; US-057584.
PR 05-SEP-1997; US-057629.
PR 05-SEP-1997; US-057642.
PR 05-SEP-1997; US-057645.
PR 05-SEP-1997; US-057648.
PR 05-SEP-1997; US-057651.
PR 05-SEP-1997; US-057662.
PR 05-SEP-1997; US-057668.
PR 05-SEP-1997; US-057762.
PR 05-SEP-1997; US-057765.
PR 05-SEP-1997; US-057771.
PR 05-SEP-1997; US-057776.
PR 06-JUN-1997; US-048876.
PR 06-JUN-1997; US-048880.
PR 06-JUN-1997; US-048883.
PR 06-JUN-1997; US-048892.
PR 06-JUN-1997; US-048895.
PR 06-JUN-1997; US-048898.
PR 06-JUN-1997; US-048901.
PR 06-JUN-1997; US-048917.
PR 06-JUN-1997; US-048963.
PR 06-JUN-1997; US-048971.
PR 06-JUN-1997; US-049019.
PR 06-JUN-1997; US-049374.
PR 05-SEP-1997; US-057627.
PR 05-SEP-1997; US-057634.
PR 05-SEP-1997; US-057643.
PR 05-SEP-1997; US-057646.
PR 05-SEP-1997; US-057649.
PR 05-SEP-1997; US-057654.
PR 05-SEP-1997; US-057654.
PR 05-SEP-1997; US-057666.
PR 05-SEP-1997; US-057760.
PR 05-SEP-1997; US-057763.
PR 05-SEP-1997; US-057769.
PR 05-SEP-1997; US-057774.
PR 05-SEP-1997; US-057777.
PR (HUMA-) HUMAN GENOME SCI INC.
PI Brewer LA, Carter KC, Dillon PJ, Ebner R, Endress GA,
PI Fan P, Feng P, Ferlie AM, Fischer CL, Florence C,
PI Florence K, Greene JM, Hu J, Kyaw H, Lafleur DW,
PI Li Y, Moore PA, Ni J, Olsen HS, Rosen CA, Ruben SM,
PI Shi Y, Soppet DR, Wei Y, Young P, Yu G, Zeng Z;
DR WPI: 99-059865/05.

DR P-PSDB: W88683, W89013, W89014, W89015, W89016.
PT New isolated human genes and the secreted polypeptides they encode -
PT useful for diagnosis and treatment of e.g. cancers, neurological
PT disorders, immune diseases, inflammation or blood disorders
PS Claim 4; Page 412-413; 772pp; English.
CC The invention relates to nucleic acid sequences (V84411 to V84633)
CC encoding human secreted proteins (W88534 to W88756). The secreted protein
CC gene sequences are deposited with the ATCC under deposit numbers ATCC
CC 97979, 97974, 97975, 97976, 97977, 209007, 209008, 209009, 209010,
CC 209011, 209080, 209081, 209082, 209083, 209084, 209085, 209511. Host
CC cells comprising recombinant vectors containing the nucleic acid
CC sequences are used for the recombinant production of the secreted
CC proteins. The polynucleotide and amino acid sequences are useful for are
CC useful for preventing, treating or ameliorating medical conditions e.g.
CC by protein or gene therapy. Pathological conditions can be also diagnosed
CC by determining the amount of the new polypeptides in a sample or by
CC determining the presence of mutations in the new polynucleotides.
CC Specific uses are described for each of the polynucleotides, based on
CC which tissues they are most highly expressed in, and include developing
CC products for the diagnosis or treatment of cancer, neurodegenerative
CC disorders, developmental abnormalities and foetal deficiencies, blood
CC disorders, tumours, leukemias, diseases of the immune system, autoimmune
CC diseases, hepatic and renal disease, lymphomas, inflammation, allergies,
CC ischemic shock, Alzheimer's and cognitive disorders, schizophrenia,
CC restenosis, prostate diseases, obesity, disorders involving osteoclasts
CC such as osteoporosis, arthritis or malignancies, diseases of testes, lung
CC or thymus, digestive/endocrine disorders, infections and AIDS. The
CC polypeptides are also useful for identifying their binding partners.
CC The present sequence represents a gene encoding a human secreted protein
CC (see descriptor line for gene number and clone identification).
SQ Sequence 1842 BP; 604 A; 322 C; 369 G; 533 T;

alignment_scores:
Quality: 35.00 Length: 8
Ratio: 4.375 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 75.000

alignment_block:
US-08-653-294-11 x V84560 ..

Align seg 1/1 to: V84560 from: 1 to: 1842

1 TyrArgLeuLeuIleArgLeuAsn 8
|||||
1560 TACAGATTACTACTACGAATGAAT 1583

seq_name: N_Geneseq_36:X20531

seq_documentation_block:
ID X20531 standard; DNA; 3858 BP.
AC X20531;
DT 05-MAY-1999 (first entry)
DE Polynucleotide sequence from the genome of Treponema pallidum.
KW Treponema pallidum infection; syphilis; Borrelia infection; animal;
KW enzyme production; ds.
OS Treponema pallidum.
PN WQ9859034-A2.
PD 30-DEC-1998.
PF 23-JUN-1998; U13041.
PR 24-JUN-1997; US-050667.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Fraser CW;
DR WPI: 99-081273/07.
PT New isolated Treponema pallidum nucleic acids - used to develop
PT products for the detection, diagnosis, characterisation, prevention
PT and therapy of T. pallidum infections, particularly syphilis
PS Claim 1; Page 368-370; 1150pp; English.
CC X20500-21243 represent polynucleotide sequences from the genome of
CC Treponema pallidum. The sequences can be used for detection,
CC diagnosis, characterisation, prevention and therapy for T. pallidum
CC infections, particularly syphilis. They can also be used for detecting
CC diseases related to Borrelia infections in animals, and for the

CC production of biosynthetic products such as enzymes. 1116 T;
SQ Sequence 3858 BP; 659 A; 777 C; 1306 G;

alignment_scores:
Quality: 35.00 Length: 9
Ratio: 4.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 77.778

alignment_block:

US-08-653-294-11 x X20531/rev ..

Align seg 1/1 to reverse of: X20531 from: 1 to: 3858

2 ArgLeuLeuIleArgLeuAsnGluArg 10

||||: ||||||||||||||||

2281 CGGATCACCATTAGCTGAACGACGCC 2255

seq_name: N_Geneseq_36:V65691

seq_documentation_block:

ID V65691 standard; DNA; 5243 BP.

AC V65691;

DT 25-JAN-1999 (first entry)

DE Fowlpox Virus vector DNA fragment.

KW Vector; poxvirus; infection; treatment; prevention; copy number;

KW promoter; vaccine; ds.

OS Fowlpox virus.

PN WC9844093-A1.

PD 08-OCT-1998:

PF 26-MAR-1998; J01358.

PR 28-MAR-1997; JP-094875.

PA (JAPG) NIPPON ZEON KK.

PI Sato T;

DI WPI: 98-557105/47.

PT Fowl poxvirus-originated 5 kb DNA vector with heterologous genes

PT useful as a vaccine, providing large copy number per cell, with

PT superior preventive efficacy

PS Claim 2; Page 36-39; 58pp; Japanese.

CC This sequence is used in the construction of a new vector originated

CC from fowlpox virus which can duplicate poxvirus in infected cells. The

CC DNA vector can be used to produce vaccines for preventing and/or treating

CC infection caused by poxvirus including orthopoxvirus and chicken

CC poxvirus. The DNA vector is about 5 kb and is concomitant. It can grow in

CC non-poxvirus-infective cells. It has a large copy number per cell, making

CC the vaccine very effective.

SQ Sequence 5243 BP; 1826 A; 882 C; 1026 G; 1508 T;

alignment_scores:

Quality: 35.00 Length: 9
Ratio: 4.375 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 66.667

alignment_block:

US-08-653-294-11 x V65691 ..

Align seg 1/1 to: V65691 from: 1 to: 5243

1 TyrArgLeuLeuIleArgLeuAsnGlu 9

|||||||:|||||

3138 TATCGACTACTCGTCAATACACGAA 3164

seq_name: N_Geneseq_36:T20013

seq_documentation_block:

ID T20013 standard; cDNA to mRNA; 307 BP.

AC T20013;

DT 17-JUL-1996 (first entry)

DE Human gene signature HUMGS01151.

KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;

KW human; cloning; mapping; non-biased library; diagnosis; detection;

KW cell typing; abnormal cell function; ss.

OS Homo sapiens.
PN W09514772-A1.
PD 01-JUN-1995.
PF 11-NOV-1994; J01916.
PR 12-NOV-1993; JP-355504.
PA (MATS/) MATSUBARA K.
PA (OKUB/) OKUBO K.
PI Matsubara K, Okubo K;
DI WPI: 95-206931/27.
PT Identifying gene signatures in 3'-directed human cDNA library - e.g.
PT for diagnosis of abnormal cell function, by preparing cDNA that
PT reflects relative abundance of corresp. mRNA in specific human
PT tissues

PS Claim 1; Pages 536-537; 2245pp; Japanese.

CC A single-stranded DNA (or its complementary strand or the corresp.

CC double-stranded DNA) which comprises one of the 7837 "GS" sequences

CC given in 119001-T26837 and which is able to hybridize to part of

CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)

CC sequences were obtained from 3'-directed cDNA libraries prepared

CC from various human tissues; synthesis of cDNA was initiated from the

CC 3'-end of mRNA by using poly(T) as the sole primer. Since the 3'-

CC untranslated sequence is unique to a particular mRNA species, almost

CC all the 3'-oriented cDNAs hybridize with specific mRNAs. Each library

CC is constructed so as to reflect accurately the relative abundance of

CC different mRNAs in the particular tissue from which it was derived.

CC The appearance frequency of a given GS in a cDNA library can be

CC determined (esp. using primers and probes derived from the GS

CC sequences) as a means of diagnosing abnormal cell function or for

CC recognising different cell types.

SQ Sequence 307 BP; 93 A; 49 C; 64 G; 101 T;

alignment_scores:

Quality: 34.00 Length: 9
Ratio: 3.778 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 55.556

alignment_block:

US-08-653-294-11 x T20013 ..

Align seg 1/1 to: T20013 from: 1 to: 307

1 TyrArgLeuLeuIleArgLeuAsnGlu 9

|||||||:|||||

258 TACAGAATATTAGTAAGATTATCA 284

seq_name: N_Geneseq_36:V77268

seq_documentation_block:

ID V77268 standard; DNA; 394 BP.

AC V77268;

DT 16-MAR-1999 (first entry)

DE Staphylococcus aureus contig SEQ ID #2957.

KW Computer readable medium; vaccine; S aureus infection; immunodetection;

KW cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;

KW skin infection; surgical wound infection; scalded skin syndrome;

KW toxic shock syndrome; ds.

OS Staphylococcus aureus.

PN EP-786519-A2.

PD 30-JUL-1997.

PF 07-JAN-1997; 100117.

PR 05-JAN-1996; US-009861.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA,

PI Rosen CA;

DI WPI: 97-374922/35.

PT Polynucleotide(s) and proteins derived from Staphylococcus aureus

PT stored on computer readable medium and used in the production of

PT anti-S.aureus vaccines

PS Claim 1; Page 2383; 3271pp; English.

CC This sequence represents one of 5191 Staphylococcus aureus DNA sequences

CC of the invention. The DNA sequences are recorded on a computer readable

CC medium, preferably selected from a floppy or hard disk, random access

CC memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
 CC the S.aureus DNA sequences allows putative functions to be assigned so
 CC that protein-encoding or regulatory regions of commercial, therapeutic or
 CC industrial importance can be obtained. Specifically, sequences which are
 CC likely to encode antigens have been identified and these polypeptides can
 CC be used in a vaccine composition against S.aureus infection. The
 CC polypeptides can also be used in a kit for the immunodetection of
 CC S.aureus in a sample. S.aureus is implicated in numerous human diseases,
 CC including cellulitis, eyelid infections, food poisoning, osteomyelitis,
 CC skin and surgical wound infections, scalded skin syndrome, toxic shock
 CC syndrome, etc. Organisms transformed with the DNA sequences can be used
 CC for recombinant production of the polypeptides. The new DNA sequences
 CC (and their fragments) are useful as primers or probes for isolating
 CC homologues of any of the S.aureus DNA sequences contained on the
 CC computer readable medium.
 SQ Sequence 394 BP; 109 A; 85 C; 71 G; 126 T;

alignment_scores:
 Quality: 34.00 Length: 10
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 60.000

alignment_block:
 US-08-653-294-11 x V77268 ..
 Align seg 1/1 to: V77268 from: 1 to: 394

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
 88 TACAATACATCGACTTTCGAGG 117

seq_name: N_Geneseq_36:X18185

seq_documentation_block:
 ID X18185 standard; DNA; 810 BP.
 AC X18185;
 DT 07-MAY-1999 (first entry)
 DE S. aureus gidB coding sequence.
 KW GidB; mutation detection; bacteriostatic; bacteriocidal compound;
 KW microbial infection; osteomyelitis; septic arthritis; gene therapy;
 KW septic thrombophlebitis; acute bacterial endocarditis; bacteraemia;
 KW cancer; ds.
 OS Staphylococcus aureus.
 PN EP-892055-A2.
 PD 20-JAN-1999.
 PF 30-JUN-1998; 305175.
 PR 12-JUN-1998; US-097072.
 PR 01-JUL-1997; US-886638.
 PA (SMIK) SMITHKLINE BEECHAM CORP.
 PI Burnham MKR, Kallender H, Palmer LM, Ward J;
 DR WPI: 99-083572/08.
 DR P-PSDB; W74405.
 PT New Staphylococcus aureus GidB polypeptides and polynucleotides -
 PT useful as diagnostic reagents and for prevention and treatment of
 PT Staphylococcus infections
 PS Claim 2; Page 21; 39pp; English.
 CC This sequence encodes the Staphylococcus aureus gidB protein of the
 CC invention. GidB polynucleotides and polypeptides are useful for
 CC diagnosing susceptibility to diseases by detecting mutations or
 CC polymorphisms in the gidB gene or analysing for the presence of amount of
 CC GidB polypeptide expressed in a patient sample. GidB PCR probes are
 CC useful for diagnosing diseases, and can characterise the response of the
 CC infectious organism to drugs. GidB polypeptides and polynucleotides are
 CC also useful for screening for antagonists, agonists and drugs against
 CC infectious micro-organisms. GidB agonists and antagonists are
 CC bacteriostatic and bacteriocidal compounds which can be used in treatment
 CC to enhance or block GidB activity, therefore treating diseases caused by
 CC microbial infection, especially S. aureus diseases including
 CC osteomyelitis, septic arthritis, septic thrombophlebitis, acute bacterial
 CC endocarditis and bacteraemia in cancer patients. Epitopes of GidB
 CC polypeptides and polynucleotides are useful immunogens for producing

CC anti-GidB antibodies for prevention of bacterial infections, and GidB
 CC polynucleotides can be used in genetic immunisation to prevent
 CC infections. GidB polypeptides, polynucleotides and their (ant)agonists
 CC can prevent adhesion of bacteria to matrix proteins, and are useful for
 CC use on wounds and body implants to prevent bacterial infection. GidB
 CC polypeptides and polynucleotides may also be used as reagents for
 CC differential screening methods.
 SQ Sequence 810 BP; 295 A; 113 C; 158 G; 244 T;

alignment_scores:
 Quality: 34.00 Length: 10
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 60.000

alignment_block:
 US-08-653-294-11 x X18185 ..

Align seg 1/1 to: X18185 from: 1 to: 810

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
 138 TATCGTTTACTTGTGATGCAATGAAAG 167

seq_name: N_Geneseq_36:X18186

seq_documentation_block:
 ID X18186 standard; DNA; 900 BP.
 AC X18186;
 DT 07-MAY-1999 (first entry)
 DE S. aureus gidB ORF sequence.
 KW GidB; mutation detection; bacteriostatic; bacteriocidal compound;
 KW microbial infection; osteomyelitis; septic arthritis; gene therapy;
 KW septic thrombophlebitis; acute bacterial endocarditis; bacteraemia;
 KW cancer; ds.
 OS Staphylococcus aureus.
 PN EP-892055-A2.
 PD 20-JAN-1999.
 PF 30-JUN-1998; 305175.
 PR 12-JUN-1998; US-097072.
 PR 01-JUL-1997; US-886638.
 PA (SMIK) SMITHKLINE BEECHAM CORP.
 PI Burnham MKR, Kallender H, Palmer LM, Ward J;
 DR WPI: 99-083572/08.
 DR P-PSDB; W74406.
 PT New Staphylococcus aureus GidB polypeptides and polynucleotides -
 PT useful as diagnostic reagents and for prevention and treatment of
 PT Staphylococcus infections
 PS Claim 2; Page 23; 39pp; English.
 CC This sequence encodes the Staphylococcus aureus gidB protein of the
 CC invention. GidB polynucleotides and polypeptides are useful for
 CC diagnosing susceptibility to diseases by detecting mutations or
 CC polymorphisms in the gidB gene or analysing for the presence of amount of
 CC GidB polypeptide expressed in a patient sample. GidB PCR probes are
 CC useful for diagnosing diseases, and can characterise the response of the
 CC infectious organism to drugs. GidB polypeptides and polynucleotides are
 CC also useful for screening for antagonists, agonists and drugs against
 CC infectious micro-organisms. GidB agonists and antagonists are
 CC bacteriostatic and bacteriocidal compounds which can be used in treatment
 CC to enhance or block GidB activity, therefore treating diseases caused by
 CC microbial infection, especially S. aureus diseases including
 CC osteomyelitis, septic arthritis, septic thrombophlebitis, acute bacterial
 CC endocarditis and bacteraemia in cancer patients. Epitopes of GidB
 CC polypeptides and polynucleotides are useful immunogens for producing
 CC anti-GidB antibodies for prevention of bacterial infections, and GidB
 CC polynucleotides can be used in genetic immunisation to prevent
 CC infections. GidB polypeptides, polynucleotides and their (ant)agonists
 CC can prevent adhesion of bacteria to matrix proteins, and are useful for
 CC use on wounds and body implants to prevent bacterial infection. GidB
 CC polypeptides and polynucleotides may also be used as reagents for
 CC differential screening methods.
 SQ Sequence 900 BP; 338 A; 124 C; 165 G; 267 T;

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alignment_scores:
  Quality: 34.00      Length: 10
  Ratio: 3.778       Gaps: 0
  Percent Similarity: 90.000  Percent Identity: 60.000

alignment_block:
  US-08-653-294-11 x X18186 ..
  Align seg 1/1 to: X18186 from: 1 to: 900

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10
  |||||
156 TATCGTTTACTTGTGTAATGAAAG 185

seq_name: N_Geneseq_36:V82078

seq_documentation_block:
  ID V82078 standard; DNA; 1300 BP.
  AC V82078;
  DE 25-MAR-1999 (first entry)
  KW GidA1; DNA encoding a partial gidA1 protein.
  KW GidA1; bacterial infection; meningitis; Helicobacter pylori infection;
  KW cancer; ulcer; gastritis; antibacterial; in-dwelling device;
  KW wound treatment; bacterial adhesion; matrix protein; ds.
  OS Staphylococcus aureus.
  FH Key Location/Qualifiers
  FT CDS
  FT 2..1165
  FT /*tag= a
  FT /transl_except= (pos: 848..850, aa: Xaa)
  FT /product= gidA1
  FT /note= "Xaa= unspecified amino acid"

EP-889129-A2.
07-JAN-1999.
30-JUN-1998; 305180.
01-JUL-1997; US-052758.
PA (SMK ) SMITHLINE BEECHAM CORP.
PA (SMK ) SMITHLINE BEECHAM PLC.
PI Burham M, Kallender H, Lenox AL, Palmer LM;
DR WPI: 99-062660/05.
DR P-PSDB: W89446.
PT New isolated gidA1 polypeptide from Staphylococcus aureus - used to
PT diagnose, treat and prevent bacterial infections e.g. S. aureus and
PT H. pylori and associated cancers, ulcers and gastritis
PS Claim 2: Page 5-6; 43pp; English.
CC The present sequence encodes a partial gidA1 protein of Staphylococcus
CC aureus. GidA1 proteins, nucleic acids and agonists are used to
CC treat conditions requiring increased activity or expression of gidA1,
CC while conditions (particularly bacterial infections) requiring
CC inhibition of gidA1 are treated by administering an antagonist,
CC inhibitory nucleic acid or competitive polypeptide. The products are
CC also used to treat S. pneumoniae infection, particularly meningitis and
CC also Helicobacter pylori infections e.g. related cancers, ulcers and
CC gastritis. These antibacterial agents may also be used to treat
CC in-dwelling devices to prevent infection or generally as wound
CC treatments to prevent adhesion of bacteria to matrix proteins.
SQ Sequence 1300 BP; 477 A; 195 C; 262 G; 365 T;

alignment_scores:
  Quality: 34.00      Length: 10
  Ratio: 3.778       Gaps: 0
  Percent Similarity: 90.000  Percent Identity: 60.000

alignment_block:
  US-08-653-294-11 x V82078 ..
  Align seg 1/1 to: V82078 from: 1 to: 1300

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10
  |||||
1249 TATCGTTTACTTGTGTAATGAAAG 1278

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seq_name: N_Geneseq_36:V74973

seq_documentation_block:
  ID V74973 standard; DNA; 1398 BP.
  AC V74973;
  DT 16-MAR-1999 (first entry)
  DE Staphylococcus aureus contig SEQ ID #662.
  KW Computer readable medium; vaccine; S.aureus infection; immunodetection;
  KW cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;
  KW skin infection; surgical wound infection; scalded skin syndrome;
  KW toxic shock syndrome; ds.
  OS Staphylococcus aureus.
  FH Key Location/Qualifiers
  FT misc_feature
  FT 1261..1320
  FT /*tag= a
  FT /note= "these bases represent a line of missing text in
  FT the sequence listing in the specification. They
  FT are included to maintain the nucleotide numbering
  FT given in the specification for this DNA sequence."

EP-786519-A2.
30-JUL-1997.
07-JAN-1997; 100117.
05-JAN-1996; US-009861.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA,
PI Rosen CA;
DR WPI: 97-374922/35.
PT Polynucleotide(s) and proteins derived from Staphylococcus aureus -
PT stored on computer readable medium and used in the production of
PT anti-S.aureus vaccines
PS Claim 1: Page 1586-1587; 3271pp; English.
CC This sequence represents one of 5191 Staphylococcus aureus DNA sequences
CC of the invention. The DNA sequences are recorded on a computer readable
CC medium, preferably selected from a floppy or hard disk, random access
CC memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
CC the S.aureus DNA sequences allows putative functions to be assigned so
CC that protein-encoding or regulatory regions of commercial, therapeutic or
CC industrial importance can be obtained. Specifically, sequences which are
CC likely to encode antigens have been identified and these polypeptides can
CC be used in a vaccine composition against S.aureus infection. The
CC polypeptides can also be used in a kit for the immunodetection of
CC S.aureus in a sample. S.aureus is implicated in numerous human diseases,
CC including cellulitis, eyelid infections, food poisoning, osteomyelitis,
CC skin and surgical wound infections, scalded skin syndrome, toxic shock
CC syndrome, etc. Organisms transformed with the DNA sequences can be used
CC for recombinant production of the polypeptides. The new DNA sequences
CC (and their fragments) are useful as primers or probes for isolating
CC homologues of any of the S.aureus DNA sequences contained on the
CC computer readable medium.
SQ Sequence 1398 BP; 461 A; 197 C; 186 G; 494 T;

alignment_scores:
  Quality: 34.00      Length: 10
  Ratio: 3.778       Gaps: 0
  Percent Similarity: 90.000  Percent Identity: 50.000

alignment_block:
  US-08-653-294-11 x V74973/rev ..
  Align seg 1/1 to reverse of: V74973 from: 1 to: 1398

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10
  |||||
1064 TATAGAGATTGTCGATTAAATCAGCGC 1035

seq_name: N_Geneseq_36:Q20995

seq_documentation_block:
  ID Q20995 standard; DNA; 1560 BP.
  AC Q20995;
  DT 19-MAY-1992 (first entry)

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DE  EH-4 gC gene.
KW  Equine herpes virus-4; glycoprotein gC; antigenic; vaccine;
KW  alphaherpesvirus; respiratory disease; cellular attachment;
KW  pathogenic; ss.
OS  Equine herpesvirus-4.
FH  Key      Location/Qualifiers
FT  cds      52..1509
FT          /*tag= a
FT          /product= EH-4_gC
PN  WO9201057-A.
PD  23-JAN-1992.
PF  04-JUL-1991; G01091.
PR  06-JUL-1990; GB-014950.
PA  (UNIU ) UNIV OF GLASGOW.
PA  (EQU1-) EQUINE VIROLOGY RES FOUN.
PI  Nicolson L, Onions DE;
DR  WPI: 92-056872/07.
P-PSDB: R20796.
DR  Nucleic acid sequence encoding EH-4 gH or gC protein - used to
PT  produce a vaccine for protection of horses against EH-4
PT  infection
PS  Claim 1: Page 23; 29pp; English.
CC  Equine dermal cells (NBL-6) were infected with EH-4 strain 1942
CC  viral DNA, purified and a BamHI library constructed in pUC9.
CC  Calcium shocked E. coli DHr cells were transformed with the
CC  recombinant plasmids. Additional clones were derived from a
CC  restriction digest of pUC9 contg. the BamHI G fragment. The
CC  nucleotide sequence of a region of BamHI G fragment spanning the gC
CC  gene was determined. by analysis of overlapping sequences (SEQ ID no
CC  2). Vaccines can be prepd. using this sequence, and they may be used
CC  to protect horses against EH-4 infection, inducing a higher level
CC  of immunity and less side-effects than other live virus vaccines.
CC  See also Q20994.
SQ  Sequence 1560 BP; 438 A; 409 C; 334 G; 379 T;

alignment_scores:
    Quality: 34.00      Length: 10
    Ratio: 3.778        Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x Q20995  ..
Align seg 1/1 to: Q20995 from: 1 to: 1560

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
|||||  |||||  |||||  |||||  |||||  |||||
409 TACAGACTAGAAATTCACCTAAACGAGCGC 438

seq_name: N_Geneseq_36:V82077

seq_documentation_block:
ID  V82077 standard; DNA; 2132 BP.
AC  V82077;
DT  25-MAR-1999 (first entry)
DE  DNA encoding a gIdAl protein.
KW  GIdAl; bacterial infection; meningitis; Helicobacter pylori infection;
KW  cancer; ulcer; gastritis; antibacterial; in-dwelling device;
KW  wound treatment; bacterial adhesion; matrix protein; ds.
OS  Staphylococcus aureus.
FH  Key      Location/Qualifiers
FT  cds      109..1986
FT          /*tag= a
FT          /product= gIdAl
PN  EP-889129-A2.
PD  07-JAN-1999.
PF  30-JUN-1998; 305180.
PR  01-JUL-1997; US-052758.
PA  (SMIK ) SMITHKLINE BEECHAM CORP.
PA  (SMIK ) SMITHKLINE BEECHAM PLC.
PI  Burnham N, Kallender H, Lenox AL, Palmer LM;
DR  WPI: 99-062660/06.

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DR  P-PSDB: W89445.
PT  New isolated gIdAl polypeptide from Staphylococcus aureus - used to
PT  diagnose, treat and prevent bacterial infections e.g. S. aureus and
PT  H. pylori and associated cancers, ulcers and gastritis
PS  Claim 2; Page 3-4; 43pp; English.
CC  The present sequence encodes a gIdAl protein of Staphylococcus
CC  aureus. GIdAl proteins, nucleic acids and agonists are used to
CC  treat conditions requiring increased activity or expression of gIdAl,
CC  while conditions (particularly bacterial infections) requiring
CC  inhibition of gIdAl are treated by administering an antagonist.
CC  inhibitory nucleic acid or competitive polypeptide. The products are
CC  used to treat S. pneumoniae infection, particularly meningitis and
CC  also Helicobacter pylori infections e.g. related cancers, ulcers and
CC  gastritis. These antibacterial agents may also be used to treat
CC  in-dwelling devices to prevent infection or generally as wound
CC  treatments to prevent adhesion of bacteria to matrix proteins.
SQ  Sequence 2132 BP; 791 A; 311 C; 430 G; 500 T;

alignment_scores:
    Quality: 34.00      Length: 10
    Ratio: 3.778        Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 60.000

alignment_block:
US-08-653-294-11 x V82077  ..
Align seg 1/1 to: V82077 from: 1 to: 2132

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
|||||  |||||  |||||  |||||  |||||  |||||
2070 TATCGTTTACTTGTGTGAATGGAATGAAAG 2099

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OM of: US-08-653-294-11 to: EST:* out_format : pfs

Date: Feb 8, 2000 4:02 AM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

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-MINMATCH=0.100 -LOOPCL=0.000 -LOPEXT=0.000 -CGAPOP=4.500
-CGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
-FGAPEXT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DELOP=6.000
-DELEXT=7.000 -START=1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -LOCALIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
-OUTFMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=1000000 -USER=US08653294
-NCPU=6 -ICPU=3 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-08-653-294-11
Query length: 10
Database: EST:*
Database sequences: 4538634
Database length: 1887831982
Search time (sec): 8553.360000

score_list:

Sequence	Strd Orig	ZScore	Escore	Len	Documentation
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gb_gss4:AO735952	+	39.00	137.15	578	HS_2261.A1.F09.T7C CIT
gb_gss1:CN5002P7	-	39.00	132.96	983	AO735952 Drosophila melanogaster
gb_gss10:AO157379	-	38.00	132.96	668	AO157379 nbx0005118r CUGI Rice
gb_gss11:AO271001	-	38.00	129.38	946	AO271001 nbx0005118r CUGI Rice
gb_gss1:CN5006G7E	-	38.00	128.18	1101	AO71730 Drosophila melanogaster
gb_gss1:CN5006G6	+	37.00	132.13	399	AO84468 Arabidopsis thaliana
gb_gss8:AO604329	+	37.00	132.23	403	AO604329 HS_2200.A1.MF.E07 CIT
gb_est22:AO1008780	+	37.00	134.57	405	AO1008780 EST203231 Normalized
gb_gss14:AO583741	+	37.00	129.75	552	AO583741 RPI-11-434G5 TV RPI-1
gb_est35:AO1820472	+	36.00	131.99	254	AO1820472 MEM00422.FOR Egg stage
gb_est10:AO185851	+	36.00	130.97	304	AO185851 SW3ICA1954SK Bruglia ma
gb_est17:AO604578	+	36.00	130.05	325	AO604578 no76b01.s1 NCI_CGAP_Ki
gb_est32:AO1758190	+	36.00	129.76	337	AO1758190 ty70e05.x1 NCI_CGAP_Ki
gb_est36:AO188192	+	36.00	129.24	360	AO188192 AV188192 Yuji Kohara
gb_gss9:AO128402	+	36.00	128.83	379	AO128402 HS_3085.A1.E07 MR CIT
gb_est26:AO1350192	+	36.00	128.01	421	AO1350192 qoz8h01.x1 NCI_CGAP_Lu
gb_gss4:AO675126	+	36.00	127.78	433	AO675126 HS_2162.A1.D10.T7C CIT
gb_gss8:AO054233	+	36.00	127.75	435	AO054233 CIT-HSP-234120.TR CIT
gb_est31:AO32239	+	36.00	127.66	440	F32239 HSPD24851 HM3 Homo sapie
gb_gss3:AO57891	+	36.00	127.50	449	B57891 CIT-HSP-2010015.TR CIT-H
gb_gss6:AO886481	+	36.00	127.50	449	AO886481 HS_5543.B2.A12.SP6E RH
gb_est16:AO557396	+	36.00	127.39	455	AO557396 n181h06.s1 NCI_CGAP_Br
gb_gss9:AO149335	+	36.00	127.39	455	AO149335 HS_2248.A2.B10.MR CIT
gb_gss10:AO215203	+	36.00	127.39	455	AO215203 HS_2190.A1.H08.MR CIT
gb_gss10:AO216648	+	36.00	127.24	464	AO216648 HS_3113.B2.E07 T7 CIT
gb_gss10:AO186906	+	36.00	126.60	503	AO186906 HS_3113.B2.E07 T7 CIT
gb_gss4:AO685122	+	36.00	126.43	514	AO685122 HS_2160.A1.F05.T7C CIT
gb_est33:AO1772130	+	36.00	126.31	522	AO1772130 EST25330 tomato resis
gb_est33:AO1779338	+	36.00	125.41	585	AO1779338 EST260217 tomato suscep
gb_est34:AO031017	+	36.00	124.68	642	AO031017 AO031017 Rice cDNA fr
gb_est30:AO165844	+	36.00	124.40	665	AO165842 AEMTBM58 Aedes aegypti
gb_gss1:CN500881	+	36.00	124.27	676	AO165844 AEMTBM57 Aedes aegypti
gb_est21:AO911731	+	36.00	122.44	852	AO151558 Drosophila melanogaster
gb_est23:AO128565	+	35.00	133.90	122	AO111731 oil5e06.s1 NCI_CGAP_G
gb_gss1:CN500522	+	35.00	132.43	147	AO128565 qa61b01.s1 Soares_feta
gb_est1:F02347	-	35.00	130.24	194	AO87848 Arabidopsis thaliana g
gb_est8:AO26266	-	35.00	129.72	207	F02347 HSCOWG062 normalized ind
gb_est4:AO294974	+	35.00	129.07	225	AO26266 ze97f02.s1 Soares_feta
gb_est35:AO186895	+	35.00	128.96	228	AV294974 AV294974 RIKEN full-le
gb_est18:AA729926	-	35.00	128.52	241	AO186895 wnl2e12.x1 NCI_CGAP_G
	-	35.00	127.43	277	AA729926 nx40f01.s1 NCI_CGAP_G

gb_est20:AA864515 - 35.00 127.20 255.29 285 | AA864515 oh55b03.s1 NCI_CGAP
gb_est24:AI233666 + 35.00 127.15 257.10 287 | AI233666 EST230354 Normalize
gb_est25:AI246027 - 35.00 126.98 262.54 293 | AI246027 qk44g11.x1 NCI_CGAP

seq_name: gb_est40:AV242923

seq_documentation_block:
LOCUS AV242923 246 bp mRNA EST 04-NOV-1999
DEFINITION RIKEN full-length enriched, 0 day neonate head Mus
musculus cDNA clone 4831414015 3' similar to NM_004576 Homo sapiens
protein phosphatase 2 (formerly 2A), regulatory subunit B (PR 52),
beta isoform (PPP2R2B) mRNA, mRNA, mRNA sequence.

ACCESSION AV242923
VERSION AV242923.1 GI:6230332
KEYWORDS house mouse.
SOURCE house mouse.
ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE
AUTHORS

Konno,H., Aizawa,K., Akahira,S., Akiyama,J., Carninci,P., Endo,T.,
Fukuda,S., Fukunishi,Y., Hara,A., Hayatsu,N., Hirozane,T., Hori,F.,
Ishii,Y., Ishikawa,T., Itoh,M., Izawa,M., Kadota,K., Kagawa,I.,
Kai,C., Kawai,J., Kikuchi,N., Kojima,Y., Koya,S., Kusakabe,M.,
Matsuyama,T., Miki,R., Mizuno,Y., Nakamura,M., Oda,H., Okazaki,Y.,
Owa,C., Ozawa,Y., Saito,H., Sano,M., Sato,K., Shibata,Y.,
Shibata,Y., Shigemoto,Y., Shiraki,T., Sogabe,Y., Sugahara,Y.,
Suzuki,H., Suzuki,H., Takahashi,F., Tateno,M., Tomimaga,N.,
Tsunoda,Y., Watabiki,A., Watanabe,S., Yamamura,T., Yasunishi,A.,
Yokota,T., Yoshiki,A., Yoshino,M., Muramatsu,M. and Hayashizaki,Y.
Riken Mouse ESTs (Konno,H., et al.)
Unpublished (1999)

TITLE
JOURNAL

COMMENT
On May 18, 1998 this sequence version replaced gi:3137751.
Contact: Yoshihide Hayashizaki
Genome Exploration Research Group, Life Science Tsukuba Center,
Genome Science Laboratory
The Institute of Physical and Chemical Research (RIKEN), Genomic
Sciences Center

3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan
Tel: +81-298-36-9013
Fax: +81-298-36-9098

Email: genome-res@rtr.riken.go.jp,
URL:http://genome.rtr.riken.go.jp/
Sasaki,N., Izawa,M., Watabiki,M., Ozawa,K., Tanaka,T., Yoneda,Y.,
Matsura,S., Carninci,P., Muramatsu,M., Okazaki,Y. and
Hayashizaki,Y.

Transcriptional sequencing: A method for DNA sequencing using RNA
polymerase. Proc. Natl. Acad. Sci. U.S.A. 95 (7), 3455-3460 (1998)
Itoh,M., Kitzunai,T., Akiyama,J., Shibata,K., Izawa,M., Kawai,J.,
Tomaru,Y., Carninci,P., Shibata,Y., Ozawa,Y., Muramatsu,M.,
Okazaki,Y. and Hayashizaki,Y.

Automated filtration-based high-throughput plasmid preparation
system. Genome Res. 9 (5), 463-470 (1999)

Carninci,P. and Hayashizaki,Y.
High-efficiency full-length cDNA cloning. Methods Enzymol. 303,
19-44 (1999)

Please visit our web site (<http://genome.rtr.riken.go.jp>) for
further details.

FEATURES
source

Location/Qualifiers
1..246
/organism="Mus musculus"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="4831414015"
/clone_lib="RIKEN full-length enriched, 0 day neonate
head"
/sex="mixed"
/tissue_type="head"
/dev_stage="0 day neonate"
/lab_host="DH10B"

/note="Site_1: Sall; Site_2: BamHI; CDNA library was
prepared and sequenced in Mouse Genome Encyclopedia
Project of Genome Exploration Research Group in Riken

```

/ssex="male"
/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coli DH10B"
BASE COUNT 207 a 97 c 90 g 180 t 4 others
ORIGIN

alignment_scores:
    Quality: 39.00      Length: 10
    Ratio: 3.900       Gaps: 0
    Percent Similarity: 100.000    Percent Identity: 60.000

```

```

Align seg 1/1 to: A0735952 from: 1 to: 578

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
|||||
94 TACAGTAAATCATGAGGATTATGACAGA 123

seq_name: gb_gss1:CNS002P7

seq_documentation_block:
LOCUS      CNS002P7          983 bp      DNA
DEFINITION Drosophila melanogaster genome survey sequence sp6 end of BAC
            BACN02A09 of DrosBAC library from Drosophila melanogaster (fruit

```

ACCESSION	AL097957	genomic survey sequence.
VERSION	AL097957.1	
KEYWORDS	GI:5609568	
SOURCE	GSS.	
ORGANISM	fruit fly.	
	Drosophila melanogaster	
	Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;	
	Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;	
	Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.	
REFERENCE	1 (bases 1 to 983)	
AUTHORS	Genoscope.	
TITLE	Direct Submission	
JOURNAL	Submitted (23-JUL-1999) Genoscope - Centre National de Sequençage	
	Bp 191 91006 Evry cedex - FRANCE (E-mail : segref@genoscope.cns.fr)	

```

- Web : www.genoscope.cns.fr)
Determination of this BAC-end sequence was carried out as part of
collaboration with the European Drosophila genome Project (EDGPP)
http://www.edgpp.ebi.ac.uk -. This Drosophila melanogaster BAC
library (Dros BAC) was made by Alain Billaud at CEPH (Centre
d'Etude du Polymorphisme Humain) with funding provided by a MRC
project grant. The DNA was prepared from embryos by Alain Buchet
and Genevieve Payan. It has been constructed in the vector
pBelobAC11.

FEATURES             Location/Qualifiers
     1..983
         /organism="Drosophila melanogaster"
         /plasmid="pBelobAC11"
         /db_xref="taxon:7227"
         /clone_lib="DrosBAC"
         /clone="BACN02A09"
         /note="end : SP6"
BASE COUNT          313 a   152 c   178 g   276 t   64 others
ORIGIN

alignment_scores:
    Quality:      39.00
    Ratio:        3.900
    Gaps:         0
Percent Similarity: 100.000  Percent Identity: 70.000

```

alignment_block:
US-08-653-294-11 x CNS002P7/rev ..
Align seg 1/1 to reverse of: CNS002P7 from: 1 to: 983
1 TyrArgLeuLeuIleArgLeuAsnGluArg 10

```

|||||:|||||:|||||:|||||:|||||:|||||
626 TATAAACTCTGATCAGATCAATAGCCGA 597

seq_name: gb_gss10:AQ157379

seq_documentation_block: 668 bp DNA GSS 12-SEP-1998
LOCUS AQ157379
DEFINITION nbxb0009118r CUGI Rice BAC Library Oryza sativa genomic clone
            nbxb0009118r, genomic survey sequence.
ACCESSION AQ157379
VERSION   AQ157379.1 GI:3592495
KEYWORDS  GSS.
SOURCE    Oryza sativa.
          ORGANISM
            Oryza sativa.
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
            Poaceae; Oryza.
REFERENCE  1 (bases 1 to 668)
AUTHORS   Wing, R.A. and Dean, R.A.
TITLE     A BAC End Sequencing Framework to Sequence the Rice Genome
JOURNAL   Unpublished (1998)
COMMENT   On Sep 10, 1998 this sequence version replaced gi:3554404.
          Contact: Wing RA
          Clemson University Genomics Institute
          100 Jordan Hall, Clemson, SC 29634, USA
          Tel: 864 656 7288
          Fax: 864 656 4293
          Email: rwing@clemson.edu
          Seq primer: GGAACAGCTAGACCGG
          Class: BAC ends
          High quality sequence start: 2
          High quality sequence stop: 389.
          High quality sequence stop: 389.

FEATURES             Location/Qualifiers
     source           1..668
                     /organism="Oryza sativa"
                     /strain="Japonica"
                     /cultivar="Nipponbare"
                     /db_xref="taxon:4530"
                     /clone="nbxb0009118r"
                     /clone_lib="CUGI Rice BAC Library"
                     /tissue_type="Leaf"
                     /lab_host="E. coli DH10B"
                     /note="Vector: pBelOBAC11; Site_1: HindIII; Site_2:
                     HindIII; Rice is one of two most popular grains in the
                     world. Half of the world population especially those
                     inhabiting highly populated areas of the humid tropics
                     and subtropics, rely on rice as their primary source of
                     carbohydrate. Monocotyledonous rice is a diploid plant
                     (2n=24) with a haploid genome equivalent of 431 Mbp
                     (Arumuganathan and Earle, 1991). The relatively small
                     genome of rice, three times larger than that of
                     Arabidopsis, makes it suitable for genomic studies. In
                     order to facilitate positional cloning, physical mapping
                     and genome sequencing of rice, we have constructed a BAC
                     library from Oryza sativa, Nipponbare variety. The
                     library contains 36,864 clones with an average insert size
                     of 128.5 Kb providing 10.9 haploid genome equivalents.
                     The deep coverage allows the isolation a particular
                     sequence with a probability of 99.9 %. Two high density
                     filters, each containing 18,432 clones (doubly spotted),
                     represent the whole library for colony screening."
BASE COUNT  157 a 204 c 141 g 166 t
ORIGIN
alignment_scores:
  Quality: 38.00 Length: 10
  Ratio: 3.800 Gaps: 0
  Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x AQ157379

```

Align seg 1/1 to: AQ157379 from: 1 to: 668

```

1 TyTArGLeuLeuIleArGLeuAsnGluArG 10
|||||:|||||:|||||:|||||:|||||:|||||
526 TATCGTCTTAGTTAGACTTAATAGCCGT 555

```

seq_name: gb_gss11:AQ271001

```

seq_documentation_block: 946 bp DNA GSS 03-NOV-1998
LOCUS AQ271001
DEFINITION nbxb001511f CUGI Rice BAC Library Oryza sativa genomic clone
            nbxb001511f, genomic survey sequence.
ACCESSION AQ271001
VERSION   AQ271001.1 GI:3824316
KEYWORDS  GSS.
SOURCE    Oryza sativa.
          ORGANISM
            Oryza sativa
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
            Poaceae; Oryza.
REFERENCE  1 (bases 1 to 946)
AUTHORS   Wing, R.A. and Dean, R.A.
TITLE     A BAC End Sequencing Framework to Sequence the Rice Genome
JOURNAL   Unpublished (1998)
COMMENT   Contact: Wing RA
          Clemson University Genomics Institute
          100 Jordan Hall, Clemson, SC 29634, USA
          Tel: 864 656 7288
          Fax: 864 656 4293
          Email: rwing@clemson.edu
          Seq primer: TAATACGACTCACTATAGG
          Class: BAC ends
          High quality sequence stop: 152.
          High quality sequence stop: 152.

```

FEATURES source

```

1..946
/organism="Oryza sativa"
/strain="Japonica"
/cultivar="Nipponbare"
/db_xref="taxon:4530"
/clone="nbxb001511f"
/clone_lib="CUGI Rice BAC Library"
/tissue_type="Leaf"
/lab_host="E. coli DH10B"
/note="Vector: pBelOBAC11; Site_1: HindIII; Site_2:
HindIII; Rice is one of two most popular grains in the
world. Half of the world population especially those
inhabiting highly populated areas of the humid tropics
and subtropics, rely on rice as their primary source of
carbohydrate. Monocotyledonous rice is a diploid plant
(2n=24) with a haploid genome equivalent of 431 Mbp
(Arumuganathan and Earle, 1991). The relatively small
genome of rice, three times larger than that of
Arabidopsis, makes it suitable for genomic studies. In
order to facilitate positional cloning, physical mapping
and genome sequencing of rice, we have constructed a BAC
library from Oryza sativa, Nipponbare variety. The
library contains 36,864 clones with an average insert size
of 128.5 Kb providing 10.9 haploid genome equivalents.
The deep coverage allows the isolation a particular
sequence with a probability of 99.9 %. Two high density
filters, each containing 18,432 clones (doubly spotted),
represent the whole library for colony screening."

```

```

BASE COUNT  260 a 202 c 181 g 303 t
ORIGIN

```

```

alignment_scores:
  Quality: 38.00 Length: 10
  Ratio: 4.222 Gaps: 0
  Percent Similarity: 90.000 Percent Identity: 80.000

```

```

alignment_block:
US-08-653-294-11 x AQ271001/rev ..
Align seg 1/1 to reverse of: AQ271001 from: 1 to: 946

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10
||||:|||||  |||
732 TATCAGCTATTAAATAGATCTAAATGAGCGT 703

seq_name: gb_gss1:CNS00G7E

seq_documentation_block:
LOCUS      CNS00G7E      1101 bp      DNA      GSS      03-JUN-1999
DEFINITION Drosophila melanogaster genome survey sequence T7 end of BAC:
BACR32013 of RPCI-98 library from Drosophila melanogaster (fruit
fly), genomic survey sequence.
ACCESSION  AL017130
VERSION     AL017130.1 GI:4951570
KEYWORDS   GSS.
SOURCE      fruit fly.
ORGANISM    Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE   1 (bases 1 to 1101)
AUTHORS     Samson,D., Saurin,W., Weissenbach,J. and Quetier,F.
TITLE       Direct Submission
JOURNAL     Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage :
            BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
            - Web : www.genoscope.cns.fr)
COMMENT     Determination of this BAC-end sequence was carried out as part of a
            collaboration with the Berkeley Drosophila Genome Project (BDGP).
            The BDGP is constructing a physical map of the Drosophila
            melanogaster genome using these BACs. For further information
            please see http://www.fruitfly.org The BDGP Drosophila
            melanogaster BAC library was prepared by Kazuoto Osoegawa and
            Aaron Mamoser in Piter de Jong's laboratory in the Department of
            Cancer Genetics at the Roswell Park Cancer Institute in Buffalo,
            NY. The library is named RPCI-98 and was constructed by partial
            EcoRI digestion of Drosophila DNA provided by the BDGP from the
            isogenic strain v2: cn bw sp, the same strain used for the BDGP's
            P1 and EST libraries. A more detailed description of the library
            and how to order individual BAC clones, the entire library, or
            filters for hybridization from the BACPAC Resource Center can be
            found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES             Location/Qualifiers
     source           1..1101
                     /organism="Drosophila melanogaster"
                     /db_xref="taxon:7227"
                     /clone_lib="RPCI-98"
                     /clone="BACR32013"
                     /note="end : T7"

BASE COUNT      305 a   247 c   220 g   294 t   35 others
ORIGIN

alignment_scores:
Quality: 38.00      Length: 10
Ratio: 3.800       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x CNS00G7E/rev ..
Align seg 1/1 to reverse of: CNS00G7E from: 1 to: 1101

1 TyrArgLeuLeuLeuArgLeuAsnGluArg 10
||||:|||||  |||
925 TACCACCTCTTTTTCAGCTGACGAGAGG 896

seq_name: gb_gss1:CNS00PG6
seq_documentation_block:
LOCUS      CNS00PG6      399 bp      DNA      GSS      28-JUN-1999
DEFINITION Arabidopsis thaliana genome survey sequence T7 end of BAC F801 of
IGF library from strain Columbia of Arabidopsis thaliana, genomic
survey sequence.
ACCESSION  AL084468
VERSION     AL084468.1 GI:5285608
KEYWORDS   GSS.
SOURCE      thale cress.
ORGANISM    Arabidopsis thaliana
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons; core
            eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
            Arabidopsi.
REFERENCE   1 (bases 1 to 399)
AUTHORS     Salanoubat,M., Choisne,N., Artiguenave,F., Brottier,P., Wincker,P.,
            Samson,D., Saurin,W., Weissenbach,J. and Quetier,F.
JOURNAL     Unpublished
REFERENCE   2 (bases 1 to 399)
AUTHORS     Genoscope.
TITLE       Direct Submission
JOURNAL     Submitted (25-JUN-1999) Genoscope - Centre National de Sequencage :
            BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr)
            - Web : www.genoscope.cns.fr)
            Location/Qualifiers
     source           1..399
                     /organism="Arabidopsis thaliana"
                     /strain="Columbia"
                     /db_xref="taxon:3702"
                     /clone_lib="IGF"
                     /clone="F801"
                     /note="end : T7"

BASE COUNT      105 a   57 c   98 g   139 t
ORIGIN

alignment_scores:
Quality: 37.00      Length: 8
Ratio: 4.625       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
US-08-653-294-11 x CNS00PG6 ..
Align seg 1/1 to: CNS00PG6 from: 1 to: 399

1 TyrArgLeuLeuLeuArgLeuAsn 8
|||||  |||
207 TACCGATTGATATTAGTTGAT 230

seq_name: gb_gss8:AQ064329
seq_documentation_block:
LOCUS      AQ064329      403 bp      DNA      GSS      04-AUG-1998
DEFINITION HS_2200_AL_ME_E07 CIT Approved Human Genomic Sperm Library D Homo
sapiens genomic clone Plate-2200 Col-13 Row-I, genomic survey
sequence.
ACCESSION  AQ064329
VERSION     AQ064329.1 GI:3378867
KEYWORDS   GSS.
SOURCE      human.
ORGANISM    Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
            Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 403)
AUTHORS     Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
            Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
            Hood,L.
TITLE       Sequence-tagged connectors: A sequence approach to mapping and
            scanning the human genome
JOURNAL     Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE     99380589
COMMENT     Contact: Mahairas GG, Wallace JC, Hood L
            High Throughput Sequencing Center

```

University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 2200 row: 1 column: 13
Class: BAC ends
High quality sequence stop: 403.

FEATURES

source

1..403
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="Plate-2200 Col-13 Row-1"
/note="male"
/note="Organ: sperm; Vector: pBelOBAC11; BAC Clones in E-Coli DH10B"

BASE COUNT 100 a 88 c 82 g 132 t 1 others
ORIGIN

alignment_scores:

Quality: 37.00 Length: 10
Ratio: 3.700 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-11 x AQ064329/rev ..

Align seg 1/1 to reverse of: AQ064329 from: 1 to: 403

1 TTAAGLeuLeuLeuAArgLeuAsnGluArg 10
|||||
395 TATCGCCCTTTTATCCAAATGGTGAAGA 366

seq_name: gb_est22:AI008780

seq_documentation_block:

LOCUS AI008780 405 bp mRNA EST 25-JAN-1999
DEFINITION EST203231 Normalized rat embryo, Bento Soares Rattus sp. cDNA clone
REMB017.3, end, mRNA sequence.

ACCESSION AI008780

VERSION AI008780.1 GI:3222612

KEYWORDS

SOURCE

ORGANISM

Rattus sp.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.

REFERENCE

AUTHORS

Lee, N.H., Glodek, A., Chandrasekhar, I., Mason, T.M., Quackenbush, J.,
Kerlavage, A.R. and Adams, M.D.

TITLE

Rat Genome Project: Generation of a Rat EST (RST) Catalog & Rat

JOURNAL

COMMENT

Unpublished (1998)
On Jan 17, 1998 this sequence version replaced gi:2044445.

CONTACT

Contact: Lee, NH

ATCC

The Institute for Genomic Research
9712, Medical Center Drive, Rockville, MD 20850, USA

Tel: (301)-838-3529

Fax: (301)-838-0208

Email: nhlee@tigr.org

Seq primer: M13-21.

FEATURES

source

1..405
Location/Qualifiers
/organism="Rattus sp."
/db_xref="ATCC (inhost):2016965"
/db_xref="taxon:10118"
/clone="REMB017"
/clone_lib="Normalized rat embryo, Bento Soares"
/dev_stage="embryo 8, 12, 18 dpc"
/note="vector: pT7n3pac; Site_1: EcoRI; Site_2: NotI"

BASE COUNT 117 a 74 c 63 g 151 t
ORIGIN

alignment_scores:

Quality: 37.00 Length: 9
Ratio: 4.111 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 66.667

alignment_block:

US-08-653-294-11 x AI008780 ..

Align seg 1/1 to: AI008780 from: 1 to: 405

1 TTAAGLeuLeuLeuAArgLeuAsnGlu 9

|||||

141 TACAAATACITTTGAGGTAAATCAA 167

seq_name: gb_gss14:AQ583741

seq_documentation_block:

LOCUS AQ583741 552 bp DNA GSS 07-JUN-1999
DEFINITION RPCI-11-434G5.TV RPCI-11 Homo sapiens genomic clone RPCI-11-434G5,
genomic survey sequence.

ACCESSION AQ583741

VERSION AQ583741.1 GI:5010851

KEYWORDS

SOURCE

ORGANISM

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS

Zhao, S., Adams, M.D., Nierman, W., Malek, J., de Jong, P. and
Venter, J.C.

TITLE

JOURNAL

COMMENT

Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
Map Building
Unpublished (1997)
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeetigr.org

Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genet cs (http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html).
Seq primer: T7

Class: BAC ends.

FEATURES

source

1..552
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="GDB:7666420"
/db_xref="taxon:9606"
/clone="RPCI-11-434G5"
/clone_lib="RPCI-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
RPCI11 Human Male BAC Library"

BASE COUNT

ORIGIN

144 a 117 c 70 g 220 t 1 others

alignment_scores:

Quality: 37.00 Length: 10
Ratio: 3.700 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 60.000

alignment_block:

REFERENCE
1 (bases 1 to 325)
AUTHORS
NCI-CCGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Sep 12, 1996 this sequence version replaced gi:1405116.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550
Email: Robert_Strausberg@nih.gov
Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: Stratagene, Inc., David B. Krizman,
Ph.D.
cDNA Library Arraying: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 682 Std Error: 0.00
Seq primer: -40m13 fwd. ET from Amersham
High quality sequence stop: 217.
Location/Qualifiers
FEATURES
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/db_xref="taxon:9606"
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/clone_lib="NCI-CGAP-AA1"
/tissue_type="adrenal adenoma"
/lab_host="SOLR (kanamycin resistant)"
/note="Organ: adrenal gland; Vector: Bluescript SK-;
Site_1: EcoRI; Site_2: XhoI; Cloned unidirectionally.
Primer: Oligo df. Two pooled bulk adrenal adenomas. 5'
adaptor sequence: 5' GAATTCGGCAGG 3' 3' adaptor
sequence: 5' CTCAGTGTGTTTTTTTTTTT 3' Average insert
size: 1.6 kb."
BASE COUNT 113 a 46 c 57 g 109 t
ORIGIN

alignment_scores:
Quality: 36.00 Length: 10
Ratio: 3.600 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x AA604578 ..
Align seg 1/1 to: AA604578 from: 1 to: 325

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
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172 TACAATCTACTATAAGATTAAACACAAA 201

seq_name: gb_est32:AI758190

seq_documentation_block:
LOCUS AI758190 337 bp mRNA EST 23-JUN-1999
DEFINITION ty70e05.x1 NCI-CGAP_Kid11 Homo sapiens cDNA clone IMAGE:2284448 3',
mRNA sequence.
ACCESSION AI758190
VERSION AI758190.1 GI:5151913
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 337)
AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
Tumor Gene Index
JOURNAL Unpublished (1997)
COMMENT On Dec 20, 1995 this sequence version replaced gi:1133827.
Contact: Robert Strausberg, Ph.D.
Tel: (301) 496-1550

Email: Robert_Strausberg@nih.gov
Tissue Procurement: Christopher Moskaluk, M.D., Ph.D., Michael R.
Emmert-Buck, M.D., Ph.D.
cDNA Library Preparation: M. Bento Soares, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40UP from Gibco.
Location/Qualifiers
FEATURES
source
1..337
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/lab_host="DH10B"
/note="Organ: kidney; Vector: p7T3D-Pac (Pharmacia) with
a modified polylinker; Site_1: Not I; Site_2: Eco RI;
Plasmid DNA from the normalized library NCI-CGAP_Kid3 was
prepared, and ss circles were made in vitro. Following HAP
purification, this DNA was used as tracer in a subtractive
hybridization reaction. The driver was PCR-amplified cDNAs
from a pool of 5,000 clones made from the same library
(clones 1322376-1323911, 1456007-1456775, and
1500552-1502855). Subtraction by Bento Soares and M.
Fatima Bonaudo."
BASE COUNT 131 a 50 c 55 g 101 t
ORIGIN

alignment_scores:
Quality: 36.00 Length: 10
Ratio: 3.600 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-11 x AI758190 ..
Align seg 1/1 to: AI758190 from: 1 to: 337

1 TyrArgLeuLeuIleArgLeuAsnGluArg 10
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242 TACAATCTACTATAAGATTAAACACAAA 271

seq_name: gb_est36:AV188192

seq_documentation_block:
LOCUS AV188192 360 bp mRNA EST 22-JUL-1999
DEFINITION AV188192 Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
embryo Caenorhabditis elegans cDNA clone yk520g12 5', mRNA
sequence.
ACCESSION AV188192
VERSION AV188192.1 GI:5570175
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE 1 (bases 1 to 360)
AUTHORS Kohara,Y., Shin-I,T., Thierry-Mieg,J., Thierry-Mieg,D., Mitsuki,H.,
Nishigaki,A., Motohashi,T., Zeng,Q., Watanabe,H., Sugimoto,A.,
Sano,M., Miyata,A., Mitani,Y., Iida,K., Uesugi,H., Sugiyama,Y. and
Nomoto,H.
Expressed genes in C.elegans
JOURNAL Unpublished (1999)
COMMENT On May 18, 1998 this sequence version replaced gi:3137896.
Contact: Yuji Kohara
Gene Library Lab
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854

Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.
Location/Qualifiers
1. 360
/organism="Caenorhabditis elegans"
/strain="N2"
/db_xref="taxon:6239"
/clone="yk520g12"
/clone_lib="Yuji Kohara unpublished cDNA:Strain N2
hermaphrodite embryo"
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/dev_stage="embryo"
BASE COUNT 94 a ' 92 c 37 g 137 t
ORIGIN

alignment_scores:
Quality: 36.00 Length: 8
Ratio: 4.500 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
US-08-653-294-11 x AV188192/rev ...
Align seg 1/1 to reverse of: AV188192 from: 1 to: 360

1 TyrArgLeuLeuIleArgLeuAsn 8
|||||
186 TATAAACTTTTGATTAGATTAAAT 163

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2000, 01:29:38 ; Search time 122.56 Seconds
(without alignments)
1.933 Million cell updates/sec

Title: US-08-653-294-13
Perfect score: 49
Sequence: 1 YRLAIRLDER 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : A_Geneseq_36.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	49	100.0	10	1 W47270	Immunomodulatory p
2	44	89.8	10	1 W47266	Immunomodulatory p
3	44	89.8	10	1 W47272	Immunomodulatory p
4	44	89.8	20	1 R32909	HLA-B*2702 CTL modu
5	44	89.8	20	1 R32911	HLA-B*2702 CTL modu
6	44	89.8	20	1 R32907	HLA-B*2702 CTL modu
7	44	89.8	20	1 R32907	HLA-B*2702 CTL modu
8	44	89.8	20	1 R32907	HLA-B*2702 CTL modu
9	44	89.8	20	1 W33778	Immunomodulating d
10	44	89.8	20	1 W33779	Immunomodulating d
11	39	79.6	20	1 W33792	Peptide B2702.84-7
12	39	79.6	20	1 W47268	Immunomodulatory p
13	39	79.6	20	1 R32910	HLA-B*2702 CTL modu
14	39	79.6	20	1 R32908	HLA-B*2702 CTL modu
15	39	79.6	20	1 R32908	HLA-B*2702 CTL modu
16	39	79.6	20	1 R32908	HLA-B*2702 CTL modu
17	32	65.3	318	1 W33793	Peptide B2702.84-7
18	32	65.3	318	1 W33793	Peptide B2702.84-7
19	32	65.3	318	1 W33793	Peptide B2702.84-7
20	32	65.3	318	1 W33793	Peptide B2702.84-7
21	32	65.3	318	1 W33793	Peptide B2702.84-7
22	31	63.3	485	1 R20796	EHV-4 9C. Nucleic
23	30	61.2	3398	1 R44430	eryA region polype
24	30	61.2	3457	1 R62504	Large polypeptide
25	30	61.2	3457	1 R62504	Large polypeptide
26	29	59.2	6	1 W47263	Bacillus subtilis
27	29	59.2	6	1 W33781	Immunomodulatory p
28	29	59.2	12	1 R32909	HLA-B*2702 CTL modu
29	29	59.2	12	1 W33798	HLA-B*2702 CTL modu
30	29	59.2	12	1 W33799	Immunomodulating d
31	29	59.2	53	1 W19361	Beta 7 integrin S3
32	29	59.2	158	1 W23586	Mabinlin MBLIII from
33	29	59.2	158	1 W23588	Mabinlin MBLIII fr
34	29	59.2	219	1 W82592	Mouse Rit ras-like

35 29 59.2 376 1 R21416 Carbonic anhydrase
36 29 59.2 377 1 R21417 Chlamydomonas carb
37 29 59.2 381 1 W98786 H. pylori GPO 121
38 29 59.2 537 1 R25534 Toxoplasma gondii
39 29 59.2 546 1 W67798 Thermococcus sp. K
40 29 59.2 663 1 W09055 Nicotiana plumbagi
41 29 59.2 711 1 W55103 Streptococcus pneu
42 29 59.2 738 1 R69849 Ethylene response
43 29 59.2 738 1 R69852 Ethylene response
44 29 59.2 738 1 R69853 Ethylene response
45 29 59.2 738 1 W73121 A. thaliana ethyle

ALIGNMENTS

RESULT 1

W47270
ID W47270 standard; peptide; 10 AA.
AC W47270;
DT 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.
FH Key Location/Qualifiers
FT Misc_difference 1.10
FT Note= "at least one of the amino acids is the D-isomer"
FN W09744052-A1.
PD 27-NOV-1997.
PF 23-APR-1997; U06705.
PR 22-MAY-1996; US-651650.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C. Krensky AM;
DR WPI; 98-018220/02.
PT Novel immunomodulatory peptide-type compound - useful for inhibiting
PT Transplant rejection
PS Claim 10; Page 36; 41pp; English.
CC The present sequence is an immunomodulatory peptide, which
CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
CC in a pharmaceutical composition together with a subtherapeutic dose
CC of an immunosuppressant, to extend the period of acceptance of a
CC transplant from a major histocompatibility complex (MHC) unmatched
CC donor, i.e. to inhibit transplant rejection. It can also be used in
CC the treatment of autoimmune diseases.
CC Peptides using the D-form amino acids are more effective
CC immunomodulators than their diastereomers or enantiomers.
SQ Sequence 10 AA;

Query Match 100.0%; Score 49; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 0.00014;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YRLAIRLDER 10
Db 1 YRLAIRLDER 10

RESULT 2

W47266
ID W47266 standard; peptide; 10 AA.
AC W47266;
DT 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.
FH Key Location/Qualifiers
FT Misc_difference 1.10

FT /note= "at least one of the amino acids is the
D-isomer

PN WO9744052-A1.
PD 27-NOV-1997.
PF 23-APR-1997; U06705.
PR 22-MAY-1996; US-651650.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 98-018220/02.
PT Novel immunomodulatory peptide-type compound - useful for inhibiting
transplant rejection
PS Claim 10; Page 36; 41pp; English.
CC The present sequence is an immunomodulatory peptide, which
comprises a Class I HLA-B alpha-1 domain sequence. It can be used
in a pharmaceutical composition together with a subtherapeutic dose
of an immunosuppressant, to extend the period of acceptance of a
transplant from a major histocompatibility complex (MHC) unmatched
donor, i.e. to inhibit transplant rejection. It can also be used in
the treatment of autoimmune diseases.
CC Peptides using the D-form amino acids are more effective
immunomodulators than their diastereomers or enantiomers.
SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.0016;
Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
DB 1 YRLAIRLNER 10

RESULT 3

ID W47272 standard; peptide; 10 AA.

AC W47272; 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.

FT Key Location/Qualifiers

FT Misc_difference 1. .10
FT /note= "at least one of the amino acids is the
D-isomer

FT WO9744052-A1.

PD 27-NOV-1997.

PF 23-APR-1997; U06705.

PR 22-MAY-1996; US-651650.

PA (STRD) UNIV LELAND STANFORD JUNIOR.

PI Clayberger C, Krensky AM;

DR WPI: 98-018220/02.

PT Novel immunomodulatory peptide-type compound - useful for inhibiting
transplant rejection

PS Claim 10; Page 36; 41pp; English.

CC The present sequence is an immunomodulatory peptide, which
comprises a Class I HLA-B alpha-1 domain sequence. It can be used
in a pharmaceutical composition together with a subtherapeutic dose
of an immunosuppressant, to extend the period of acceptance of a
transplant from a major histocompatibility complex (MHC) unmatched
donor, i.e. to inhibit transplant rejection. It can also be used in
the treatment of autoimmune diseases.

CC Peptides using the D-form amino acids are more effective
immunomodulators than their diastereomers or enantiomers.

CC Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;

Best Local Similarity 90.0%; Pred. No. 0.0016;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
DB 1 YRLAIRLNER 10

RESULT 4

ID R92909 standard; peptide; 20 AA.

AC R92909;

DT 16-MAY-1996 (first entry)

DE HLA-B*2702 CTL modulating peptide (B2702.84-75/75-84(T)).

KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
immunosuppressant; graft versus host disorder; transplantation; therapy;
class I MHC; HLA-B*2702.

OS Synthetic.

PN WO9526979-A1.

PD 12-OCT-1995.

PF 05-APR-1995; U04349.

PR 05-APR-1994; US-222851.

PA (STRD) UNIV LELAND STANFORD JUNIOR.

PI Clayberger C, Krensky AM, Parham P;

DR WPI: 95-358582/46.

PT Extension of acceptance period of transplants from MHC unmatched

donor hosts - using Class I B*75-84 MHC antigen of the recipient

PT host

PS Example 15; Page 36; 80pp; English.

CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
class I major histocompatibility complex (MHC) antigens. This sequence
is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
I MHC HLA-B*2702. These sequences can be used to extend the period of
acceptance by a recipient of a transplant from an MHC unmatched donor.

CC The peptides are administered to a patient in conjunction with a
subtherapeutic amount of an immunosuppressant. This is administered to
the patient for a limited period of time (compared to the lifetime
administration for current treatments). The peptides particularly
modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
of the patient.

CC Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;

Best Local Similarity 90.0%; Pred. No. 0.0035;

Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
DB 1 YRLAIRLNER 10

RESULT 5

ID R92911 standard; peptide; 20 AA.

AC R92911;

DT 16-MAY-1996 (first entry)

DE HLA-B*2702 CTL modulating peptide (B2702.84-75/84-75).

KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
immunosuppressant; graft versus host disorder; transplantation; therapy;
class I MHC; HLA-B*2702.

OS Synthetic.

PN WO9526979-A1.

PD 12-OCT-1995.

PF 05-APR-1995; U04349.

PR 05-APR-1994; US-222851.

PA (STRD) UNIV LELAND STANFORD JUNIOR.

PI Clayberger C, Krensky AM, Parham P;

DR WPI: 95-358582/46.

PT Extension of acceptance period of transplants from MHC unmatched

donor hosts - using Class I B*75-84 MHC antigen of the recipient

PT host

PS Example 15; Page 36; 80pp; English.

CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
class I major histocompatibility complex (MHC) antigens. This sequence
is a dimer of residues 75-84 of the alpha-1 domain of the class I MHC

CC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 YRLAIRLDER 10
 | | | | | | | | | |
 Db 1 YRLAIRLNER 10
 | | | | | | | | | |
 RESULT 6
 R92907
 ID R92907 standard; peptide; 20 AA.
 AC R92907; 1996 (first entry)
 DT 16-MAY-1996
 DE HLA-B2702 CTL modulating peptide (B2702.84-75/75-84).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW Immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW Class I MHC; HLA-B2702.
 OS Synthetic.
 PN WO9526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 PT WPI: 95-358382/46.
 DR Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B7-84 MHC antigen of the recipient
 PT host
 PS Example 15; Page 36; 80pp; English.
 CC R3061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC Class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 1 YRLAIRLDER 10
 | | | | | | | | | |
 Db 1 YRLAIRLNER 10
 | | | | | | | | | |

RESULT 7
 R95428
 ID R95428 standard; peptide; 20 AA.
 AC R95428;
 DT 12-NOV-1996 (first entry)
 DE HLA-B2702 84-75-84 palindrome.
 KW HLA; p74; alpha-helix; human-leucocyte-associated antigen; inhibitor;
 KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
 KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;

KW cytolysis; antigen presenting cell.
 OS Synthetic.
 PN WO9513288-A1.
 PD 18-MAY-1995.
 PF 10-NOV-1994; U12985.
 PR 10-NOV-1993; US-150493.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI: 95-194027/25.
 DT Compns. comprising lymphoid surface membrane proteins - which may
 PT inhibit cytolytic activity and differentiation of CTLs.
 PS Example; Page 12; 29pp; English.
 CC R95413, and R94415-R95431 represent palindromes and fragments of
 CC human-leucocyte-associated antigens. This sequence represents the
 CC HLA-B2702 84-75-84 palindrome. These sequences can be used to isolate
 CC the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
 CC protein associated with T-cell activation in mammalian T-cells, and is
 CC also immunologically cross reactive with the heat shock protein Hsc70.
 CC p74 is found in a limited number of cell types, but is particularly
 CC expressed on B and T cells. p74 can be isolated by lysis of a suitable
 CC cell with an amphoteric detergent, and then passed through an affinity
 CC column containing a covalently bound HLA-B2702 palindromic peptide.
 CC Compositions comprising the extracellular fragment of p74 combined with
 CC HLA-B2702.60-84 (see R95416), induces calcium influx, and inhibits
 CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
 CC compounds can be screened for their effect on the cytolytic activity of
 CC T-cells, by combining them with the extracellular portion of p74 and
 CC determining the amount of binding between the candidate compound and p74.
 CC Modulation of CTL activity can be inhibited in a cellular composition
 CC containing T-cells and antigen presenting cells (APCs), by adding to the
 CC mix the extracellular portion of p74, in an amount sufficient to compete
 CC with p74 for the binding of the p74 ligand.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
 | | | | | | | | | |
 Db 1 YRLAIRLNER 10
 | | | | | | | | | |
 RESULT 8
 W33778
 ID W33778 standard; peptide; 20 AA.
 AC W33778;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #1.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 DT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 16; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is
 CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa'6-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa'6 = E or
 CC V; aa'7 = D, S or N; aa'9 = R or G; aa80 = I or N; aa81, aa84 = a

CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alpha domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
 |||||:|
 Db 1 YRLAIRLNER 10

RESULT 9

W33779
 ID W33779 standard; peptide: 20 AA.
 AC W33779;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #2.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 16; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is
 CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alpha domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10

Db 1 YRLAIRLNER 10
 |||||:|

RESULT 10

W33792
 ID W33792 standard; peptide: 20 AA.
 AC W33792;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.84-75/75-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W3784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match

89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10

|||||:|

Db 1 YRLAIRLNER 10

RESULT 11

W47268
 ID W47268 standard; peptide: 10 AA.
 AC W47268;
 DT 22-MAY-1998 (first entry)
 DE Immunomodulatory peptide.
 KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
 KW transplant rejection; treatment; autoimmune disease.
 OS Homo sapiens.
 OS Synthetic.

Query Match 89.8%; Score 44; DB 1; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.0035;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10

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PF 23-APR-1997; U06705.
PR 22-MAY-1996; US-651650.
PA (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 98-018220/02.
PT Novel immunomodulatory peptide-type compound - useful for inhibiting
transplant rejection
PS Claim 10; Page 36; 41pp; English.
CC The present sequence is an immunomodulatory peptide, which
comprises a Class I HLA-B alpha-1 domain sequence. It can be used
in a pharmaceutical composition together with a subtherapeutic dose
of an immunosuppressant, to extend the period of acceptance of a
transplant from a major histocompatibility complex (MHC) unmatched
donor, i.e. to inhibit transplant rejection. It can also be used in
the treatment of autoimmune diseases.
CC Peptides using the D-form amino acids are more effective
immunomodulators than their diastereomers or enantiomers.
SQ Sequence 10 AA;

Query Match 79.6%; Score 39; DB 1; Length 10;
Best Local Similarity 80.0%; Pred. No. 0.019;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
   ||| |||:|
Db 1 YRLRLRNER 10

RESULT 12
R92910
ID R92910 standard; peptide; 20 AA.
AC R92910;
DT 16-MAY-1996 (first entry)
DE HLA-B2702 CTL modulating peptide (B2702.84-75(T)/75-84(T)).
KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW Immunosuppressant; graft versus host disorder; transplantation; therapy;
KW Class I MHC; HLA-B2702.
OS Synthetic.
PN W09526979-A1.
PD 12-OCT-1995.
PF 05-APR-1995; U04349.
PR (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM, Parham P;
DR WPI: 95-358582/46.
PT Extension of acceptance period of transplants from MHC unmatched
donor hosts - using Class I B75-84 MHC antigen of the recipient
PT host
PS Example 15; Page 36; 80pp; English.
CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
Class I major histocompatibility complex (MHC) antigens. This sequence
is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
I MHC HLA-B2702. These sequences can be used to extend the period of
acceptance by a recipient of a transplant from an MHC unmatched donor.
CC The peptides are administered to a patient in conjunction with a
subtherapeutic amount of an immunosuppressant. This is administered to
the patient for a limited period of time (compared to the lifetime
administration for current treatments). The peptides particularly
modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
of the patient.
SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
Best Local Similarity 80.0%; Pred. No. 0.04;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
   ||| |||:|
Db 1 YRLATRLNER 10

RESULT 13
R92908
ID R92908 standard; peptide; 20 AA.
AC R92908;
DT 16-MAY-1996 (first entry)
DE HLA-B2702 CTL modulating peptide (B2702.84-75(T)/75-84).
KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW Immunosuppressant; graft versus host disorder; transplantation; therapy;
KW Class I MHC; HLA-B2702.
OS Synthetic.
PN W09526979-A1.
PD 12-OCT-1995.
PF 05-APR-1995; U04349.
PR (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM, Parham P;
DR WPI: 95-358582/46.
PT Extension of acceptance period of transplants from MHC unmatched
donor hosts - using Class I B75-84 MHC antigen of the recipient
PT host
PS Example 15; Page 36; 80pp; English.
CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
Class I major histocompatibility complex (MHC) antigens. This sequence
is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
I MHC HLA-B2702. These sequences can be used to extend the period of
acceptance by a recipient of a transplant from an MHC unmatched donor.
CC The peptides are administered to a patient in conjunction with a
subtherapeutic amount of an immunosuppressant. This is administered to
the patient for a limited period of time (compared to the lifetime
administration for current treatments). The peptides particularly
modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
of the patient.
SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
Best Local Similarity 80.0%; Pred. No. 0.04;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
   ||| |||:|
Db 1 YRLATRLNER 10

RESULT 14
R95430
ID R95430 standard; peptide; 20 AA.
AC R95430;
DT 12-NOV-1996 (first entry)
DE HLA-B2702 84-75T/75-84T palindrome.
KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN W09513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PI (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 95-194027/25.
PT Compens. comprising lymphoid surface membrane proteins - which may
inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 12; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
human-leucocyte-associated antigens. These sequences can be used to
isolate the protein p74 from a T-cell lysate. p74 is a T-cell surface
membrane protein associated with T-cell activation in mammalian T-cells,
and is also immunologically cross reactive with the heat shock protein
Hsc70. p74 is found in a limited number of cell types, but is
particularly expressed on B and T cells. p74 can be isolated by lysis of

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CC a suitable cell with an amphoteric detergent, and then passed through an
 CC affinity column containing a covalently bound HLA-B2702 palindromic
 CC peptide. Compositions comprising the extracellular fragment of p74
 CC combined with HLA-B2702.60-84 (see R95416), induces calcium influx, and
 CC inhibits cytotoxic T lymphocyte (CTL) differentiation or cytotoxicity.
 CC Candidate compounds can be screened for their effect on the cytolytic
 CC activity of T-cells, by combining them with the extracellular portion of
 CC p74 and determining the amount of binding between the candidate compound
 CC and p74. Modulation of CTL activity can be inhibited in a cellular
 CC composition containing T-cells and antigen presenting cells (APCs), by
 CC adding to the mix the extracellular portion of p74, in an amount
 CC sufficient to compete with p74 for the binding of the p74 ligand.
 CC Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 88.9%; Pred. No. 0.04;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9
 DB 1 YRLAIRLNE 9

RESULT 15

W33791 ID W33791 standard; peptide; 20 AA.
 AC W33791;
 DE 19-JUN-1998 (first entry)
 DT Peptide B2702.84-75T/75-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-085530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 CC Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.04;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10

DB 1 YRLATRLNER 10

Search completed: February 8, 2000, 01:29:38
 Job time: 1750 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:25 ; Search time 117.7 seconds
(without alignments)
4.008 Million cell updates/sec

Title: US-08-653-294-15
Perfect score: 49
Sequence: 1 YRLIIRLDER 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : PIR_62:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	37	75.5	151	2 C71113	probable frxA prot
2	35	71.4	506	2 S37583	RING finger protei
3	35	71.4	513	1 TVHURF	ret finger protein
4	35	71.4	801	4 TVHURF	transforming prote
5	34	69.4	87	2 D70886	hypothetical prote
6	33	67.3	132	2 S75953	hypothetical prote
7	33	67.3	469	2 A71322	hypothetical prote
8	32	65.3	52	2 T07269	hypothetical prote
9	32	65.3	341	2 D48435	cysteine proteinas
10	32	65.3	342	2 S61978	hypothetical prote
11	32	65.3	348	2 I37271	cylindrii - human
12	32	65.3	554	1 F70548	probable memb prot
13	32	65.3	1400	2 B70963	hypothetical prote
14	31	63.3	30	2 S25666	phosphopyruvate hy
15	31	63.3	157	2 H75054	hypothetical prote
16	31	63.3	221	2 S56283	hypothetical prote
17	31	63.3	237	2 H75202	biotin operon repr
18	31	63.3	395	2 T01392	leucine-rich repea
19	31	63.3	560	2 I50372	ORF2 - chicken
20	31	63.3	588	2 E75060	hydrogenase-4 comp
21	31	63.3	825	2 S54455	YTA12 protein prec
22	31	63.3	1489	2 S60416	DNA helicase YglU5
23	31	63.3	1711	1 A47392	chromodomain-helic
24	30	61.2	85	2 S03746	negative sporulati
25	30	61.2	96	2 S03219	hypothetical prote
26	30	61.2	149	2 H75047	f420-nonreducing h
27	30	61.2	156	2 A70968	hypothetical prote
28	30	61.2	173	2 S27599	hypothetical prote
29	30	61.2	198	2 E69779	transcription regu
30	30	61.2	235	2 A71236	probable biotin--[

31 30 61.2 287 2 I39689 hypothetical prote
32 30 61.2 297 2 T12615 ribosomal protein
33 30 61.2 319 2 JC6117 transcription fact
34 30 61.2 324 2 G69515 transcription regu
35 30 61.2 333 2 T05643 hypothetical prote
36 30 61.2 354 1 VGBE67 glycoprotein D pre
37 30 61.2 367 2 T02529 myb-related protei
38 30 61.2 381 2 D64525 GDP-D-mannose dehy
39 30 61.2 391 2 E72539 hypothetical prote
40 30 61.2 404 2 F64238 hypothetical prote
41 30 61.2 447 2 T16527 hypothetical prote
42 30 61.2 456 2 JC6523 26s proteasom p55
43 30 61.2 513 1 I58311 HMG-box containing
44 30 61.2 540 2 T12704 leucine-rich prote
45 30 61.2 595 2 JQ1684 anthranilate synth

ALIGNMENTS

RESULT 1

C71113
Probable frxA protein - Pyrococcus horikoshii
C:Species: Pyrococcus horikoshii
C:Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 14-Aug-1998
C:Accession: C71113
R:Kawarabayashi, Y.; Sawada, M.; Horikawa, H.; Hino, Y.; Yamanoto, S.; Ogu
M.; Ohfuku, Y.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; Kushida, N.; Ogu
DNA Res. 5, 55-76, 1998
A:Title: Complete sequence and gene organization of the genome of a hyper-thermophilic
A:Reference number: A71000; MUID:98344137
A:Accession: C71113
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-151 <RAW>
A:Cross-references: GB:AP000003; MID:g3236130; PID:d1030708; PID:g3257082
A:Experimental source: strain OT3
A:Note: This accession replaces an interim accession for a sequence replaced by GenBa
C:Genetics:
A:Gene: PH0674

Query Match 75.5%; Score 37; DB 2; Length 151;
Best Local Similarity 88.9%; Pred. No. 1.8;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLLIRLDER 10
|||||
DB 96 RLLIELDER 104

RESULT 2

S37583
RING finger protein rfp - mouse (fragment)
C:Species: Mus musculus (house mouse)
C:Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 16-Jul-1999
C:Accession: S37583
R:Takahashi, M.
submitted to the EMBL Data Library, October 1993
A:Reference number: S37583
A:Accession: S37583
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-506 <RAW>
A:Cross-references: EMBL:X75343; MID:9406747; PIDN:CAA53092.1; PID:g406748
C:Superfamily: rfp transforming protein; RING finger homology
C:Keywords: zinc
F:5-55/Domain: RING finger homology <RNG>

Query Match 71.4%; Score 35; DB 2; Length 506;
Best Local Similarity 77.8%; Pred. No. 1.6;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDE 9
|||||
Db 191 YRLARLEE 199

RESULT 3
TVHURF
ret finger protein - human
N:Alternate names: transforming protein rfp
C:Species: Homo sapiens (man)
C:Date: 30-Sep-1989 #sequence_revision 30-Sep-1989 #text_change 18-Jun-1999
C:Accession: A28101
R:Takahashi, M.; Inaguma, Y.; Hiai, H.; Hirose, F.
Mol. Cell. Biol. 8, 1853-1856, 1988
A:Title: Developmentally regulated expression of a human "finger"-containing gene encode
A:Reference number: A28101; MUID:88246464
A:Accession: A28101
A:Molecule type: mRNA
A:Residues: 1-513 <TAK>
A:CROSS-references: DDBJ:J03407; NID:g337371; PIDN:AAA36564.1; PID:g337372
C:Genetics:
A:Gene: GDB:RFP
A:CROSS-references: GDB:511359; GDB:1391662
A:Map position: 6p22-6p21.3
C:Superfamily: rfp transforming protein; RING finger homology
C:Keywords: DNA binding; transforming protein; zinc
F:1-315/Product: transforming protein rfp (fragment) #status predicted <RET>
F:12-62/Domain: RING finger homology <RNG>
F:16-127/Domain: metal and nucleic acid binding #status predicted <TMN>

Query Match 71.4%; Score 35; DB 1; Length 513;
Best Local Similarity 77.8%; Pred. No. 16;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDE 9
|||||
Db 198 YRLARLEE 206

RESULT 4
TVHURF
transforming protein RFP/protein-tyrosine kinase RET mutant fusion protein - human
N:Alternate names: ret oncogene protein
N:Contains: protein-tyrosine kinase (EC 2.7.1.112) ret
C:Species: Homo sapiens (man)
C:Date: 31-Mar-1989 #sequence_revision 10-Sep-1997 #text_change 13-Aug-1999
C:Accession: A27203
R:Takahashi, M.; Cooper, G.M.
Mol. Cell. Biol. 7, 1378-1385, 1987
A:Title: ret transforming gene encodes a fusion protein homologous to tyrosine kinases.
A:Reference number: A27203; MUID:87257826
A:Accession: A27203
A:Molecule type: mRNA
A:Residues: 'QAGA', 1-801 <TAK>
A:CROSS-references: GB:M16029; NID:g340025
A:Note: codons preceding the probable start codon were translated
C:Comment: The ret oncogene is the chimeric product of a translocation mutation between
C:Genetics:
A:Gene: RFP/RET
C:Keywords: ATP; fusion protein; oncogene; phosphotransferase; transforming protein; ty
F:1-315/Region: transforming protein rfp
F:316-792/Region: protein-tyrosine kinase ret
F:459-467/Region: protein kinase ATP-binding motif
F:487/Active site: Lys #status predicted

Query Match 71.4%; Score 35; DB 4; Length 801;
Best Local Similarity 77.8%; Pred. No. 26;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDE 9

Db 198 YRLARLEE 206
|||||
RESULT 5
D70886
hypothetical protein Rv2866 - Mycobacterium tuberculosis (strain H37RV)
C:Species: Mycobacterium tuberculosis
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 17-Jul-1998
C:Accession: D70886
R:Cole, S.T.; Davies, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
R:Conor, R.; Brown, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd,
Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
Nature 393, 537-544, 1998
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno
A:Reference number: A70500; MUID:98295987
A:Accession: D70886
A>Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-87 <COL>
A:CROSS-references: GB:AL008883; GB:AL123456; NID:g3261490; PID:ell72958; PID:g261281
A:Experimental source: strain H37RV
C:Genetics:
A:Gene: Rv2866

Query Match 69.4%; Score 34; DB 2; Length 87;
Best Local Similarity 66.7%; Pred. No. 4.2;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDE 9
|||||
Db 60 YRLIRIDD.68

RESULT 6
S75953
hypothetical protein - Synecocystis sp. (strain PCC 6803)
C:Species: Synecocystis sp.
A:Variety: PCC 6803
C:Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 21-Aug-1998
C:Accession: S75953
R:Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima,
O. K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yas
DNA Res. 3, 109-136, 1996
A:Title: Sequence analysis of the genome of the unicellular cyanobacterium Synecocys
s.
A:Reference number: S74322; MUID:97061201
A:Accession: S75953
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-132 <KAN>
A:CROSS-references: EMBL:D64006; GB:AB001339; NID:gl001291; PID:dl011451; PID:gl00131
A:Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996

Query Match 67.3%; Score 33; DB 2; Length 132;
Best Local Similarity 66.7%; Pred. No. 10;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDE 9
|||||
Db 120 YRLIRLDD 128

RESULT 7
A71322
hypothetical protein TP0471 - syphilis spirochete
C:Species: Treponema pallidum subsp. pallidum (syphilis spirochete)
C:Date: 24-Jul-1998 #sequence_revision 24-Jul-1998 #text_change 17-Mar-1999
C:Accession: A71322
R:Fraser, C.M.; Norris, S.J.; Weinstock, G.M.; White, O.; Sutton, G.G.; Dodson, R.; G

rson, J.; Khalak, H.; Richardson, D.; Howell, J.K.; Chidambaram, M.; Utterback, T.; McDevitt, L.; Weidman, J.; Smith, H.O.; Venter, J.C.
 Science 281, 375-388, 1998
 A:Title: Complete genome sequence of *Treponema pallidum*, the syphilis spirochete.
 A:Reference number: A71250; MUID:98332770
 A:Accession: A71322
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-469 <COL>
 A:Cross-references: GB:AE001223; GB:AE000520; NID:g3322745; PID:g3322760
 A:Experimental source: strain Nichols
 C:Genetics:
 A:Gene: TP0471

Query Match 67.3%; Score 33; DB 2; Length 469;
 Best Local Similarity 87.5%; Pred. No. 38;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIIRLD 8
 | | | | |
 Db 140 YTLIIRLD 147

RESULT 8

T07269
 hypothetical protein 52b - *Chlorella vulgaris* chloroplast
 C:Species: Chloroplast *Chlorella vulgaris*
 C:Date: 14-May-1999 #sequence_revision 14-May-1999 #text_change 22-Jun-1999
 C:Accession: T07269
 R:Wakasugi, T.; Nagai, T.; Kapoor, M.; Sugita, M.; Ito, M.; S.; Tsudzuki, J.; Nakas
 Proc. Natl. Acad. Sci. U.S.A. 94, 5967-5972, 1997
 A:Title: Complete nucleotide sequence of the chloroplast genome from the green alga *Chlorella vulgaris*
 A:Reference number: Z15985; MUID:97303241
 A:Accession: T07269
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-52 <WAK>
 A:Cross-references: EMBL:AB001684; NID:d1110444; PID:d1021495
 C:Genetics:
 A:Genome: chloroplast
 C:Keywords: chloroplast

Query Match 65.3%; Score 32; DB 2; Length 52;
 Best Local Similarity 60.0%; Pred. No. 6.4;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLIIRLD 10
 : | | | | |
 Db 17 FLIIIVELDER 26

RESULT 9

D48435
 cysteine proteinase AC-3 - nematode (*Haemonchus contortus*)
 C:Species: *Haemonchus contortus*
 C:Date: 01-Dec-1993 #sequence_revision 18-Nov-1994 #text_change 22-Jun-1999
 C:Accession: D48435
 R:Pratt, D.; Ames, L.G.; Hageman, R.; Reynolds, V.; Boisvenue, R.J.; Cox, G.N.
 Mol. Biochem. Parasitol. 51, 209-218, 1992
 A:Title: Cloning and sequence comparisons of four distinct cysteine proteases expressed
 A:Reference number: A48435; MUID:92244291
 A:Accession: D48435
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-341 <PRA>
 A:Cross-references: GB:M80388; NID:g159178; PID:AAA29178.1; PID:g159179
 A:Note: sequence extracted from NCBI backbone (NCBIN:98512, NCBI:P:98520)
 C:Superfamily: papain

Query Match 65.3%; Score 32; DB 2; Length 341;

Best Local Similarity 60.0%; Pred. No. 44;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLIIRLD 10
 | | | | |
 Db 221 YRLIIRMDKR 230

RESULT 10
 S61978
 hypothetical protein YPL088w - yeast (*Saccharomyces cerevisiae*)
 N:Alternate names: hypothetical protein YPL088w
 C:Species: *Saccharomyces cerevisiae*
 C:Date: 10-Apr-1996 #sequence_revision 19-Apr-1996 #text_change 29-Sep-1999
 C:Accession: S61978
 R:Wang, Y.; Ahmed, A.; Bussey, H.; Fortin, N.; Friesen, J.D.; Hall, J.; Storms, R.K.;
 submitted to the EMBL Data Library, December 1995
 A:Description: The sequence of *Saccharomyces cerevisiae* chromosome XVI left arm.
 A:Reference number: S61959
 A:Accession: S61978
 A:Molecule type: DNA
 A:Residues: 1-342 <WAN>
 A:Cross-references: EMBL:U43281; NID:g1151218; PIDN:AA68211.1; PID:g1151238; MIPS:YP
 C:Genetics:
 A:Map position: 16L
 C:Superfamily: conserved hypothetical protein YPL088w

Query Match 65.3%; Score 32; DB 2; Length 342;
 Best Local Similarity 70.0%; Pred. No. 45;
 Matches 7; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLIIRLD 10
 | | | | |
 Db 207 YNLIYREDER 216

RESULT 11

I37271
 cylicin II - human
 C:Species: *Homo sapiens* (man)
 C:Date: 12-Aug-1996 #sequence_revision 12-Aug-1996 #text_change 25-Oct-1996
 C:Accession: I37271; S52774
 R:Hess, H.; Heid, H.; Zimbelmann, R.; Franke, W.W.
 Exp. Cell Res. 218, 174-182, 1995
 A:Title: The protein complexity of the cytoskeleton of bovine and human sperm heads:
 A:Reference number: I37271; MUID:95255491
 A:Accession: I37271
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-348 <HES>
 A:Cross-references: EMBL:Z46788; NID:g758586; PID:g758587

Query Match 65.3%; Score 32; DB 2; Length 348;
 Best Local Similarity 60.0%; Pred. No. 45;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLIIRLD 10
 | | | | |
 Db 78 YRLIIRISER 87

RESULT 12

F70548
 probable menD protein - *Mycobacterium tuberculosis* (strain H37RV)
 C:Species: *Mycobacterium tuberculosis*
 C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999
 C:Accession: F70548
 R:Coile, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
 ; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd,
 Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.
 Nature 393, 537-544, 1998

A:Authors: Sqaers, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome
A:Reference number: A70500; MUID:98295987

A:Accession: F70548

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-554 <COL>

A:Cross-references: GB:295558; GB:AL123456; NID:g3261781; PID:e316800; PID:g2114017

A:Experimental source: strain H37RV

C:Genetics:

A:Gene: mend

C:Superfamily: menD protein

Query Match 65.3%; Score 32; DB 1; Length 554;

Best Local Similarity 66.7%; Pred. No. 73;

Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLRLRDLR 10

||:||||

DB 48 RLHVRIDER 56

RESULT 13

B70963

hypothetical protein Rv0236c - Mycobacterium tuberculosis (strain H37RV)

C:Species: Mycobacterium tuberculosis

C>Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 17-Jul-1998

C:Accession: B70963

R: Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.

; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd, S.

Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.

Nature 393, 537-544, 1998

A:Authors: Sqaers, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome

A:Reference number: A70500; MUID:98295987

A:Accession: B70963

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-1400 <COL>

A:Cross-references: GB:292669; GB:AL123456; NID:g3242271; PID:e1300719; PID:g3242273

A:Experimental source: strain H37RV

C:Genetics:

A:Gene: Rv0236c

Query Match 65.3%; Score 32; DB 2; Length 1400;

Best Local Similarity 75.0%; Pred. No. 1.9e+02;

Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 3 LLRLRDLR 10

||:||||

DB 621 VLLRLDER 628

RESULT 14

S25666

phosphopyruvate hydratase (EC 4.2.1.11) - Streptomyces hygroscopicus (fragment)

C:Species: Streptomyces hygroscopicus

C>Date: 04-Dec-1992 #sequence_revision 04-Dec-1992 #text_change 18-Jul-1997

C:Accession: S25666

R: Hidaka, T.; Hidaka, M.; Uozumi, T.; Seto, H.

Mol. Gen. Genet. 233, 476-478, 1992

A:Title: Nucleotide sequence of a carboxyphosphoenolpyruvate phosphonmutase gene isol

ns.

A:Reference number: S23585; MUID:92318902

A:Accession: S25666

A:Molecule type: DNA

A:Residues: 1-30 <HID>

A:Cross-references: EMBL:D00609

C:Experimental source: strain SF1293

C:Superfamily: enolase

C:Keywords: carbon-oxygen lyase; gluconeogenesis; glycolysis; hydro-lyase

Query Match 63.3%; Score 31; DB 2; Length 30;

Best Local Similarity 66.7%; Pred. No. 5.8;

Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLRLRDL 9

||:||||

DB 17 YNQLRLDE 25

RESULT 15

H75054

hypothetical protein PAB2372 - Pyrococcus abyssi (strain Orsay)

C:Species: Pyrococcus abyssi

C>Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Aug-1999

C:Accession: H75054

R: anonymous, Genoscope

submitted to the EMBL Data Library, July 1999

A:Description: Pyrococcus abyssi genome sequence: Insights into archaeal chromosome s

A:Reference number: A75001

A:Accession: H75054

A>Status: preliminary

A:Molecule type: DNA

A:Residues: 1-157 <KAW>

A:Cross-references: GB:AJ248287; GB:AL096836; NID:g5458657; PIDN:CABS0333.1; PID:e151

A:Experimental source: strain Orsay

C:Genetics:

A:Gene: PAB2372

Query Match 63.3%; Score 31; DB 2; Length 157;

Best Local Similarity 75.0%; Pred. No. 32;

Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRDL 8

||||:|

DB 137 YRLRLRDL 144

Search completed: February 7, 2000, 11:54:26

Job time: 24336 sec

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2000, 00:59:53 ; Search time 63.71 Seconds
(without alignments)
4.688 Million cell updates/sec

Title: US-08-653-294-15
Perfect score: 49
Sequence: 1 YRLRLRDLR 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 82229 seqs, 29864866 residues

Total number of hits satisfying chosen parameters: 82229

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries
Database : SwissProt_38:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	35	71.4	513	1 RFP_HUMAN	P14373 homo sapien
2	35	71.4	522	1 RFP_MOUSE	Q62158 mus musculus
3	32	65.3	299	1 RL5_BOMMO	Q76190 bombyx mori
4	32	65.3	317	1 OLF3_CANFA	Q95156 canis famli
5	32	65.3	317	1 OLF3_HUMAN	Q13607 homo sapien
6	32	65.3	348	1 CYL2_HUMAN	Q14093 homo sapien
7	31	63.3	221	1 YF88_YEAST	P43592 saccharomyc
8	31	63.3	327	1 RL5_ANOGA	O44248 anopheles g
9	31	63.3	627	1 GIDA_COXBU	P94613 coxiellia bu
10	31	63.3	825	1 RCAL_YEAST	P40341 saccharomyc
11	31	63.3	1489	1 YGPQ_YEAST	P53115 saccharomyc
12	31	63.3	1709	1 CHD1_HUMAN	O14646 homo sapien
13	31	63.3	1711	1 CHD1_MOUSE	P40201 mus musculus
14	30	61.2	85	1 SPOE_BACSU	P05043 bacillus su
15	30	61.2	96	1 YL12_SSV1	P20219 sulfolobus
16	30	61.2	294	1 RL5A_SCHPO	P52822 schizosacch
17	30	61.2	294	1 RL5B_SCHPO	O74306 schizosacch
18	30	61.2	297	1 RL5_HELAN	O65353 helianthus
19	30	61.2	354	1 VGLI_VZVD	P09258 varicella-z
20	30	61.2	404	1 Y349_MYCGE	P47591 mycoplasma
21	30	61.2	595	1 TRPE_ARATH	P32068 arabidopsis
22	30	61.2	611	1 EMPA_VIBAN	P43147 vibrio angu
23	30	61.2	621	1 GIDA_HELPY	P56138 helicobacte
24	30	61.2	621	1 TRPX_ARATH	P32069 arabidopsis
25	30	61.2	628	1 GIDA_BACSU	P25812 bacillus su
26	30	61.2	629	1 GIDA_HAEIN	P44763 haemophilus
27	30	61.2	635	1 TRG4_ECOLI	Q00185 escherichia
28	30	61.2	637	1 TRG5_ECOLI	Q00184 escherichia
29	30	61.2	659	1 YVBT_BACSU	P37484 bacillus su
30	30	61.2	851	1 STA2_HUMAN	P52630 homo sapien
31	30	61.2	864	1 STA2_PIG	O02799 sus scrofa
32	30	61.2	899	1 SECA_BORBU	O07497 borrelia bu
33	30	61.2	3079	1 IRA2_YEAST	P19158 saccharomyc
34	30	61.2	3224	1 N358_HUMAN	P49792 homo sapien

35	29	59.2	101	1 YEB4_YEAST	P39999 saccharomyc
36	29	59.2	124	1 RL5_PIG	Q95276 sus scrofa
37	29	59.2	172	1 USC2_YEAST	P06104 saccharomyc
38	29	59.2	179	1 UBC2_CANAL	O74201 candida alb
39	29	59.2	293	1 DBFB_PSEPA	P47243 pseudomonas
40	29	59.2	293	1 RL5_CAEEL	P49405 caenorhabdi
41	29	59.2	295	1 RL5A_XENLA	P15125 xenopus lae
42	29	59.2	295	1 RL5B_XENLA	P15126 xenopus lae
43	29	59.2	296	1 RL5_CHICK	P22451 gallus gall
44	29	59.2	296	1 RL5_HUMAN	P46777 homo sapien
45	29	59.2	296	1 RL5_RAT	P09895 rattus norv

ALIGNMENTS

RESULT 1
RFP_HUMAN
ID RFP_HUMAN STANDARD; PRT; 513 AA.
AC P14373;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE ZINC-FINGER PROTEIN RFP (RET FINGER PROTEIN).
GN RFP.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Homnidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 88246464.
RA TAKAHASHI M., INAGUMA Y., HIAI H., HIROSE F.;
RT "Developmentally regulated expression of a human 'finger'-containing
RT gene encoded by the 5' half of the ret transforming gene.";
RL Mol. Cell. Biol. 8:1853-1856(1988).
CC -!- FUNCTION: MAY FUNCTION IN MALE GERM CELL DEVELOPMENT.
CC -!- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC -!- DISEASE: RECOMBINATION OF THE N-TERMINAL OF RFP WITH A PROTEIN
CC TYROSINE KINASE PRODUCES THE RET TRANSFORMING PROTEIN.
CC -!- SIMILARITY: CONTAINS A C3HC4-CLASS ZINC FINGER.
CC
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CC
CC EMBL; J03407; AAA36564.1; -.
CC PIR; A28101; TVHURF.
CC MIM; 602165; -.
CC PROSITE; PS00518; ZINC_FINGER_C3HC4; 1.
CC PFAM; PF00097; ZF-C3HC4; 1.
CC PFAM; PF00622; SPRY; 1.
CC PFAM; PF00643; zf-B_box; 1.
CC Proto-oncogene; Zinc-finger; Metal-binding; Chromosomal translocation;
CC Nuclear protein; DNA-binding.
CC SITE 315 316
CC BREAKPOINT FOR TRANSLOCATION TO FORM THE
CC RFP-RET ONCOGENE.
CC ZN_FING 16 56
CC DOMAIN 96 127
CC B_BOX.
CC SEQUENCE 513 AA; 58489 MW; 022BC859 CRC32;
CC

Query Match 71.4%; Score 35; DB 1; Length 513;
Best Local Similarity 77.8%; Pred. No. 8.3;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRDLR 9

||||| |||

Db 198 YRLRLARLEE 206

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RESULT 2
RFP_MOUSE STANDARD; PRT; 522 AA.
ID QF2158;
AC 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE ZINC-FINGER PROTEIN RFP (RET FINGER PROTEIN).
GN RFP.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-C57BL/6;
RX MEDLINE: 97176437.
RA CAO T., SHANNON M., HANDEL M.A., ETKIN L.D.;
RT "Mouse ret finger protein (rfp) proto-oncogene is expressed at
RT specific stages of mouse spermatogenesis.";
RT Dev. Genet. 19:309-320(1996).
CC -1- FUNCTION: MAY FUNCTION IN MALE GERM CELL DEVELOPMENT.
CC -1- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
CC -1- SIMILARITY: CONTAINS A C3HC4-CLASS ZINC FINGER.
CC
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CC
CC EMBL; L48855; AAA8354.1;
DR MGD; MGI:97904; RFP.
DR PROSITE; PS00518; ZINC_FINGER_C3HC4; 1.
DR PFAM; PF00097; zf-C3HC4; 1.
DR PFAM; PF00622; SPRY; 1.
DR PFAM; PF00643; zf-B_box; 1.
KW Zinc-finger; Metal-binding; Nuclear protein; DNA-binding.
FT ZN_FING 25 65 C3HC4-TYPE.
FT DOMAIN 105 136 B_BOX.
SQ SEQUENCE 522 AA; 59550 MW; 18E6E716 CRC32;

Query Match 71.4%; Score 35; DB 1; Length 522;
Best Local Similarity 77.8%; Pred. No. 8.5;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 YRLIRLDE 9
DB 207 YRLARLEE 215
|||||||
-----
RESULT 3
RL5_BOMMO STANDARD; PRT; 299 AA.
ID 076190;
AC 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE 60S RIBOSOMAL PROTEIN L5.
GN RPL5.
OS Bombyx mori (Silk moth).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Lepidoptera; Glossata; Ditrysia;
OC Bombycoidea; Bombycidae; Bombyx.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-SILK GLAND;
RA YANG C.S., SEHNAL F.;
RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: THIS PROTEIN BINDS 5S RNA (BY SIMILARITY).

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CC -1- SIMILARITY: BELONGS TO THE L18P FAMILY OF RIBOSOMAL PROTEINS.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AF008229; AAC24960.1;
DR PFAM; PF00861; Ribosomal_L18p; 1.
KW Ribosomal protein; rRNA-binding.
SQ SEQUENCE 299 AA; 34378 MW; 7262D2FC CRC32;

Query Match 65.3%; Score 32; DB 1; Length 299;
Best Local Similarity 50.0%; Pred. No. 20;
Matches 5; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

OY 1 YRLIRLDER 10
DB 49 YRLIVRLSNK 58
|||||||
-----
RESULT 4
OLF3_CANFA STANDARD; PRT; 317 AA.
ID OLF3_CANFA
AC Q95136;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE OLFACTORY RECEPTOR-LIKE PROTEIN OLF3.
OS Canis familiaris (Dog).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Carnivora; Fissipedia; Canidae; Canis.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 97008103.
RA ISSEL-TARVER L., RINE J.;
RT "Organization and expression of canine olfactory receptor genes.";
RL Proc. Natl. Acad. Sci. U.S.A. 93:10897-10902(1996).
CC -1- FUNCTION: PUTATIVE ODORANT OR SPERM CELL RECEPTOR.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
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CC
CC EMBL; U53681; AAB37241.1;
DR GCRDB; GCR_1190;
DR PROSITE; PS00237; G_PROTEIN_RECEPTOR; 1.
DR PFAM; PF00001; 7tm_1; 1.
KW G-protein coupled receptor; Transmembrane; Glycoprotein;
KW Multigene family; Olfaction.
FT DOMAIN 1 25 EXTRACELLULAR (POTENTIAL).
FT TRANSNM 26 49 1 (POTENTIAL).
FT TRANSNM 50 57 CYTOPLASMIC (POTENTIAL).
FT TRANSNM 58 79 2 (POTENTIAL).
FT DOMAIN 80 100 EXTRACELLULAR (POTENTIAL).
FT TRANSNM 101 120 3 (POTENTIAL).
FT DOMAIN 121 139 CYTOPLASMIC (POTENTIAL).
FT TRANSNM 140 158 4 (POTENTIAL).
FT DOMAIN 159 196 EXTRACELLULAR (POTENTIAL).
FT TRANSNM 197 219 5 (POTENTIAL).
FT TRANSNM 220 236 CYTOPLASMIC (POTENTIAL).
FT TRANSNM 237 260 6 (POTENTIAL).
FT DOMAIN 261 272 EXTRACELLULAR (POTENTIAL).

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FT TRANSMEM 273 292 7 (POTENTIAL).
FT DOMAIN 293 317 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 5 POTENTIAL.
SQ SEQUENCE 317 AA; 35238 MW; C16156EE CRC32;

Query Match 65.3%; Score 32; DB 1; Length 317;
Best Local Similarity 87.5%; Pred. No. 21;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 LLIRLDR 10
Db 47 LLIRLDR 54

RESULT 5
OLF3_HUMAN STANDARD; PRT; 317 AA.
ID OLF3_HUMAN
AC Q13607;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE OLFACTORY RECEPTOR-LIKE PROTEIN OLF3.
GN OLF3.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.

RA ISSEL-TARVER L., RINE J.;
RL Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: PUTATIVE ODORANT RECEPTOR.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -!- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
CC
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CC
CC EMBL; U56421; AA01215.1; -
CC GCRDB; GCR1924; -
CC DR PROSITE; PS00237; G.PROTEIN_RECEPTOR; 1.
CC PFAM; PF00001; 7tm_1; 1.
CC
CC G-protein coupled receptor; Transmembrane; Glycoprotein;
CC Multigene family; Olfaction.
CC
CC DOMAIN 1 24 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 25 48 1 (POTENTIAL).
CC DOMAIN 49 57 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 58 79 2 (POTENTIAL).
CC DOMAIN 80 100 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 101 120 3 (POTENTIAL).
CC DOMAIN 121 139 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 140 160 4 (POTENTIAL).
CC DOMAIN 161 200 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 201 222 5 (POTENTIAL).
CC DOMAIN 223 236 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 237 261 6 (POTENTIAL).
CC DOMAIN 262 272 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 273 292 7 (POTENTIAL).
CC DOMAIN 293 317 CYTOPLASMIC (POTENTIAL).
CC CARBOHYD 5 POTENTIAL.
CC DISULFID 97 189 BY SIMILARITY.
CC SEQUENCE 317 AA; 35316 MW; 30A35A35 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 317;
Best Local Similarity 87.5%; Pred. No. 21;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 3 LLIRLDR 10
Db 47 LLIRLDR 54

RESULT 6
CYL2_HUMAN STANDARD; PRT; 348 AA.
ID CYL2_HUMAN
AC Q14093;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE CYLICIN II (MULTIPLE-BAND POLYPEPTIDE II).
GN CYL2.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-TESTIS;
RX MEDLINE; 95255491.
RA HESS H., HEID H., ZIMBELMANN R., FRANK W.W.;
RT "The protein complexity of the cytoskeleton of bovine and human sperm
heads: the identification and characterization of cylicin II.";
RL Exp. Cell Res. 218:174-182(1995).
CC -!- FUNCTION: POSSIBLE ARCHITECTURAL ROLE DURING SPERMATOGENESIS. MAY
CC BE INVOLVED IN SPERMATID DIFFERENTIATION.
CC -!- SUBCELLULAR LOCATION: CALYX; SPERM HEAD CYTOSKELETAL STRUCTURE.
CC -!- TISSUE SPECIFICITY: TESTIS.
CC
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CC
CC EMBL; Z46789; CAA86752.1; -
CC Cytoskeleton; Structural protein; Repeat; Sperm; Spermatogenesis.
CC FT DOMAIN 25 347 31 X 3 AA REPEATS OF K-K-X.
CC FT DOMAIN 157 240 3 X APPROXIMATE TANDEM REPEATS.
CC FT REPEAT 157 184 1.
CC FT REPEAT 185 212 2.
CC FT REPEAT 213 240 3.
CC SEQUENCE 348 AA; 39079 MW; FD27FBEF CRC32;

Query Match 65.3%; Score 32; DB 1; Length 348;
Best Local Similarity 60.0%; Pred. No. 23;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 1 YRLIRLDR 10
Db 78 YRLIRLDR 87

RESULT 7
YFH8_YEAST STANDARD; PRT; 221 AA.
ID YFH8_YEAST
AC P43592;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE HYPOTHETICAL 25.9 KD PROTEIN IN MPRI-GCN20 INTERGENIC REGION.
GN YFR008W.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomycetes.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-S288C / AB972;
RX MEDLINE; 95400292.

RA MURAKAMI Y., NAITOU M., HAGIWARA H., SHIBATA T., OZAWA M.,
 RA SASANUMA S.-I., SASANUMA M., TSUCHIYA Y., SOEDA E., YOKOYAMA K.,
 RA YAMAZAKI M., TASHIRO H., EKI T.,
 RT "Analysis of the nucleotide sequence of chromosome VI from
 RT Saccharomyces cerevisiae";
 RL Nat. Genet. 10:261-268(1995).
 CC -----
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 CC -----
 CC EMBL; D50617; BAA09247.1; -
 DR Hypothetical protein. 22
 KW DOMAIN 11 POLY-GLN.
 SQ SEQUENCE 221 AA; 25915 MW; 3D34764A CRC32;

 Query Match 63.3%; Score 31; DB 1; Length 221;
 Best Local Similarity 85.7%; Pred. No. 23;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

 QY 1 YRLRL 7
 Db 211 YRLRL 217

 RESULT 8
 ID RL5_ANOGA STANDARD; PRT; 327 AA.
 AC 04248;
 DT 15-DEC-1998 (Rel. 37, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 15-DEC-1998 (Rel. 37, Last annotation update)
 DE 60S RIBOSOMAL PROTEIN L5.
 GN RPL5.
 OS Anopheles gambiae (African malaria mosquito).
 OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
 OC Pterygota; Neoptera; Endopterygota; Diptera; Nematocera; Culicoides;
 OC Culicidae; Anophelinae.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=G3;
 RA CORNEL A.J., KUMAR V., MUKABAYIRE O., SALAZAR RAFFERTY C.,
 RA PETRARCA V., COLUZZI M., COLLINS F.H.;
 RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: THIS PROTEIN BINDS 5S RNA (BY SIMILARITY).
 CC -!- SIMILARITY: BELONGS TO THE L18P FAMILY OF RIBOSOMAL PROTEINS.
 CC -----
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 CC -----
 CC EMBL; AF002236; AAB97731.1; -
 DR PFAM; AF00861; Ribosomal_L18p; 1.
 KW Ribosomal protein; rRNA-binding.
 SQ SEQUENCE 327 AA; 37996 MW; F3A3EED2 CRC32;

 Query Match 63.3%; Score 31; DB 1; Length 327;
 Best Local Similarity 50.0%; Pred. No. 35;
 Matches 5; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

 QY 1 YRLRL 10
 Db 49 FRLVRLSNR 58

RESULT 9
 GIDA_COXBU STANDARD; PRT; 627 AA.
 ID GIDA_COXBU
 AC P94613;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 15-JUL-1998 (Rel. 36, Last annotation update)
 DE GLUCOSE INHIBITED DIVISION PROTEIN A.
 GN GIDA.
 OS Coxiella burnetii.
 OC Bacteria; Proteobacteria; gamma subdivision; Coxiella.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=NINE MILE PHASE I;
 RA WILLEMS H., JAEGER C.;
 RL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: NOT KNOWN.
 CC -!- SIMILARITY: BELONGS TO THE GIDA FAMILY.
 CC -----
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 CC -----
 CC EMBL; Y10436; CAA71459.1; -
 DR PROSITE; PS01280; GIDA_1; 1.
 DR PROSITE; PS01281; GIDA_2; 1.
 DR PFAM; PF01134; GIDA; 1.
 SQ SEQUENCE 627 AA; 69951 MW; B9AF4071 CRC32;

 Query Match 63.3%; Score 31; DB 1; Length 627;
 Best Local Similarity 75.0%; Pred. No. 70;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

 QY 1 YRLRL 8
 Db 436 YRLRL 443

 RESULT 10
 ID RCAL_YEAST STANDARD; PRT; 825 AA.
 AC P40341;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE MITOCHONDRIAL RESPIRATORY CHAIN COMPLEXES ASSEMBLY PROTEIN RCAL
 DE (EC 3.4.24.-) (FAT-BINDING HOMOLOG 12).
 GN RCAL OR YTA12 OR YMR089C OR YMR582.14C.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Saccharomyces.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC MEDLINE; 95014441.
 RA TZAGOLOFF A., YUE J., JANG J., JANG M.F.;
 RT "A new member of a family of ATPases is essential for assembly of
 RT mitochondrial respiratory chain and ATP synthetase complexes in
 RT Saccharomyces cerevisiae";
 RL J. Biol. Chem. 269:26144-26151(1994).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C;
 RX MEDLINE; 95274317.
 RA SCHNALL R., MANNHAUPT G., STUCKA R., TAUER R., EHNLE S.,
 RA SCHWARZLOSE C., VETTER I., FELDMANN H.;
 RT "Identification of a set of yeast genes coding for a novel family of

RT putative ATPases with high similarity to constituents of the 26S
 RT protease complex.
 RL Yeast 10:1141-1155(1994).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN-S288C / AB972;
 RA GENTLES S., BOWMAN S., BARRELL B.G., RAJANDREAM M.A.;
 RL Submitted (MAY-1995) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: PUTATIVE ATP-DEPENDENT PROTEASE. NECESSARY FOR THE
 CC ASSEMBLY OF MITOCHONDRIAL RESPIRATORY CHAIN AND ATPASE COMPLEXES.
 CC FUNCTION BOTH IN POSTTRANSLATIONAL ASSEMBLY AND IN THE TURNOVER OF
 CC MISTRANSLATED OR MISTOLDED POLYPEPTIDES.
 CC -!- COFACTOR: BINDS ONE ZINC ION (POTENTIAL).
 CC -!- SUBCELLULAR LOCATION: MITOCHONDRIAL.
 CC -!- SIMILARITY: BELONGS TO THE AAA FAMILY OF ATPASES.
 CC -!- SIMILARITY: BELONGS TO PEPTIDASE FAMILY M41 (ZINC
 CC METALLOPROTEASE).
 CC -----
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 CC -----
 DR EMBL; U09358; AAA62606.1; -;
 DR EMBL; X81068; CAA56955.1; -;
 DR EMBL; Z49259; CAA89236.1; -;
 DR SGD; L0002584; YTA12.
 DR PROSITE; PS00674; AAA; 1.
 DR PFAM; PF01434; Peptidase_M41; 1.
 DR KW ATP-binding; Mitochondrion; Transmembrane; Hydrolase; Metalloprotease;
 KW Zinc.
 FT TRANSMEM 178 194 POTENTIAL.
 FT TRANSMEM 294 311 POTENTIAL.
 FT NP_BIND 388 395 ATP (POTENTIAL).
 FT METAL 613 613 ZINC (CATALYTIC) (BY SIMILARITY).
 FT ACT_SITE 614 614 BY SIMILARITY.
 FT METAL 617 617 ZINC (CATALYTIC) (BY SIMILARITY).
 FT CONFLICT 349 350 DV -> EL (IN REF. 2).
 FT CONFLICT 653 653 I -> V (IN REF. 1).
 SQ SEQUENCE 825 AA; 93276 MW; 69EBD054 CRC32;

 Query Match 63.3%; Score 31; DB 1; Length 825;
 Best Local Similarity 77.8%; Pred. No. 93;
 Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 YRLRLRLE 9
 |||||
 Db 36 YRLNRLQE 44

 RESULT 11
 YGPO_YEAST
 ID YGPO_YEAST STANDARD; PRT; 1489 AA.
 AC P53115;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-NOV-1997 (Rel. 35, Last annotation update)
 DE HYPOPHOSPHATE 171.5 K D HELICASE IN NUT1-ARO2 INTERGENIC REGION.
 GN YGL150C OR G1880.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 CC Saccharomycetaceae; Saccharomycetes.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 96158061.
 RA JAMES C.M., INDE K.J., OLIVER S.G.;
 RT "DNA sequence analysis of a 35 kb segment from Saccharomyces
 RT cerevisiae chromosome VII reveals 19 open reading frames including

RT RAD54, ACE1/CUP2, PMR1, RCK1, AMS1 and CAL1/CDC43.";
 RL Yeast 11:1413-1419(1995).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-S288C / FY1769;
 RX MEDLINE; 97197983.
 RA VOT M., DEFOOR E., VERHASSELT P., RILES L., ROBBEN J., VOLCKAERT G.;
 RL "The sequence of a nearly unclonable 22.8 kb segment on the left arm
 RT chromosome VII from saccharomyces cerevisiae reveals ARO2, RPL9A,
 RT TIP1, MRP1 genes and six new open reading frames.";
 RL Yeast 13:177-182(1997).
 CC -!- SUBCELLULAR LOCATION: NUCLEAR (POTENTIAL).
 CC -!- SIMILARITY: BELONGS TO THE SNF2/RAD54 HELICASE FAMILY.
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 CC -----
 DR EMBL; Z48618; CAA88537.1; -;
 DR EMBL; Z72672; CAA96861.1; -;
 DR EMBL; X99960; CAA68224.1; -;
 DR PFAM; PF00176; SNF2_N; 1.
 DR PFAM; PF00271; helicase_C; 1.
 DR KW Hypothetical protein; Nuclear protein; DNA-binding; Helicase;
 KW ATP-binding.
 FT DOMAIN 188 193 POLY-ALA.
 FT DOMAIN 259 268 POLY-GLU.
 FT DOMAIN 300 306 POLY-SER.
 FT DOMAIN 568 573 POLY-GLU.
 FT DOMAIN 675 682 POLY-GLU.
 FT NP_BIND 731 738 ATP (POTENTIAL).
 FT SITE 841 844 DEAD BOX.
 SQ SEQUENCE 1489 AA; 171454 MW; 8149887E CRC32;

 Query Match 63.3%; Score 31; DB 1; Length 1489;
 Best Local Similarity 58.3%; Pred. No. 1; 7e+02;
 Matches 7; Conservative 3; Mismatches 0; Indels 2; Gaps 1;
 QY 1 YRLIR-LDER 10
 |||||
 Db 1420 YRLVRGTIEER 1431

 RESULT 12
 CHD1_HUMAN
 ID CHD1_HUMAN STANDARD; PRT; 1709 AA.
 AC Q14646;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE CHROMODOMAIN-HELICASE-DNA-BINDING PROTEIN 1 (CHD-1).
 GN CHD1.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 CC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 97470991.
 RA WOODAGE T., BASRAI M.A., BAXEVANIS A.D., HIETER P., COLLINS F.S.;
 RT "Characterization of the CHD family of proteins.";
 RL Proc. Natl. Acad. Sci. U.S.A. 94:11472-11477(1997).
 CC -!- FUNCTION: SEQUENCE-SELECTIVE DNA-BINDING PROTEIN. COULD PLAY AN
 CC IMPORTANT ROLE IN GENE REGULATION.
 CC -!- SUBCELLULAR LOCATION: NUCLEAR.
 CC -!- SIMILARITY: BELONGS TO THE SNF2/RAD54 HELICASE FAMILY.
 CC -!- SIMILARITY: CONTAINS 2 'CHROMO' DOMAINS.
 CC -----
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DR EMBL; Z99111; CAB13237.1; -.
DR PIR; S03746; S03746.
DR SUBTILIST; BG10769; SPOOE.
KW Sporulation; Transcription regulation.
SQ SEQUENCE 85 AA; 9791 MW; E2B23676 CRC32;

Query Match 61.2%; Score 30; DB 1; Length 85;
Best Local Similarity 55.6%; Pred. No. 14;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLLRLDER 10
| | | : | | |
Db 9 RLLVSIK 17

RESULT 15

Y112_SSV1 STANDARD; PRT; 96 AA.
ID Y112_SSV1
AC P20219;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 01-AUG-1992 (Rel. 23, Last annotation update)
DE HYPOTHETICAL 11.2 KD PROTEIN (ORF E-96).
OS Sulfolobus virus-like particle SSV1.
OC Viruses; dsDNA viruses, no RNA stage; Fuselloviridae; Fusellovirus.
RN (1)
RP SEQUENCE FROM N.A.
RX MEDLINE; 92024080.
RA PALM P., SCHLEPER C., GRAMPP B., YEATS S., MCWILLIAM P., REITER W.-D.,
ZILLIG W.;
RT "Complete nucleotide sequence of the virus SSV1 of the
archaeobacterium Sulfolobus shibatae.";
RL Virology 185:242-250(1991).
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CC -----
DR EMBL; X07234; CAA30218.1; -.
DR PIR; S03219; S03219.
KW Hypothetical protein.
SQ SEQUENCE 96 AA; 11176 MW; D351EB9B CRC32;

Query Match 61.2%; Score 30; DB 1; Length 96;
Best Local Similarity 44.4%; Pred. No. 16;
Matches 4; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 YRLRLDE 9
| : : : | | |
Db 75 YKILRCDE 83

Search completed: February 8, 2000, 00:59:54
Job time: 3783 sec

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Sequence	Strd	Orig	ZScore	EScore	Len	Documentation
gb_gss11: AQ324186	-	39.00	142.06	37.98	784	I AQ324186 mgb00017A14r CUGI Rice
gb_est5: D75467	-	37.00	141.45	41.05	335	I D75467 CELK104H2F Yuij Kohara unp
gb_est5: D73508	+	37.00	140.81	44.57	360	I D73508 CELK051H2F Yuij Kohara unp
gb_est5: D73590	+	37.00	140.81	44.57	360	I D73590 CELK060G3F Yuij Kohara unp
gb_est5: D74728	+	37.00	140.81	44.57	360	I D74728 CELK084F3F Yuij Kohara unp
gb_est5: D75423	+	37.00	140.81	44.57	360	I D75423 CELK104C7F Yuij Kohara unp
gb_est5: D75727	+	37.00	140.81	44.57	360	I D75727 CELK109D2F Yuij Kohara unp
gb_est16: C40575	+	37.00	140.81	44.57	360	I C40575 C40575 Yuij Kohara unp
gb_est16: C42212	+	37.00	140.81	44.57	360	I C42212 C42212 Yuij Kohara unp
gb_est16: C44981	+	37.00	140.81	44.57	360	I C44981 C44981 Yuij Kohara unp
gb_est16: C51139	+	37.00	140.81	44.57	360	I C51139 C51139 Yuij Kohara unp
gb_est36: AV192603	+	37.00	140.81	44.57	360	I AV192603 AV192603 Yuij Kohara unp
gb_est36: AV192774	+	37.00	140.81	44.57	360	I AV192774 AV192774 Yuij Kohara unp
gb_est36: AV193922	+	37.00	140.81	44.57	360	I AV193922 AV193922 Yuij Kohara unp
gb_est36: AV194609	+	37.00	140.81	44.57	360	I AV194609 AV194609 Yuij Kohara unp
gb_est36: AV191482	+	37.00	140.57	45.99	370	I AV191482 AV191482 Yuij Kohara unp
gb_est16: C39480	+	37.00	140.47	46.56	374	I C39480 C39480 Yuij Kohara unp
gb_est16: C42649	+	37.00	140.47	46.56	374	I C42649 C42649 Yuij Kohara unp
gb_est16: C48686	+	37.00	140.42	46.84	375	I C48686 C48686 Yuij Kohara unp
gb_est16: C43167	+	37.00	140.40	46.98	377	I C43167 C43167 Yuij Kohara unp
gb_est36: AV191244	+	37.00	140.37	47.13	378	I AV191244 AV191244 Yuij Kohara unp
gb_est28: A1531870	+	37.00	138.72	58.25	455	I A1531870 SD03153.Sprime SD CRO
gb_gss8: AQ308578	-	37.00	138.38	60.90	473	I AQ308578 CIT-HSP-2333K12.TR CIT
gb_est12: AA280470	+	36.00	136.67	73.90	354	I AA280470 SWFCA2019SK Brugia ma
gb_gss9: A0146935	-	36.00	136.63	98.47	455	I A0146935 HS_2248_A2.B10.MR CIT
gb_gss8: AQ038143	-	35.00	132.77	124.93	354	I AQ038143 CIT-HSP-2320M23.TR CIT
gb_gss12: AQ331915	-	35.00	132.26	133.22	372	I AQ331915 HS_5002_A2.D11.SP6E RP
gb_est36: AV189109	+	35.00	132.26	133.22	375	I AV189109 AV189109 Yuij Kohara unp
gb_est17: C73120	-	35.00	132.44	168.55	460	I C73120 C73120 Rice panicle at f
gb_est23: A0021869	-	35.00	129.73	184.56	498	I A0021869 A0021869 Mouse unfert
gb_gss11: AQ294280	+	35.00	129.71	184.98	501	I AQ294280 HS_3000_B1.F08.MR CIT
gb_gss14: AQ543593	+	35.00	129.68	185.83	499	I AQ543593 RPCI-11-359C3.TV RPCI-1
gb_est44: AV390501	+	35.00	128.89	205.46	547	I AV390501 AV390501 Chlamydomonas
gb_est34: A0068481	-	34.00	132.00	137.99	244	I A0068481 A0068481 Rice callus C
gb_est44: A0088183	-	34.00	130.01	178.10	305	I A0088183 A0081183 Oncorhynchus
gb_est41: D36382	+	34.00	129.31	194.89	330	I D36382 CELK032EF Yuij Kohara unp
gb_gss11: AQ281049	+	34.00	128.38	208.45	350	I AQ281049 RPCI11-76J9.TJ RPCI-11
gb_est2: T98600	-	34.00	128.68	211.18	354	I T98600 ye60e09.sl Soares fetal
gb_est1: D36749	-	34.00	128.53	215.27	360	I D36749 CELK036F4F Yuij Kohara unp
gb_est36: AV201748	-	34.00	128.53	215.27	360	I AV201748 AV201748 Yuij Kohara unp
gb_est17: C63035	-	34.00	128.19	224.87	374	I C63035 C63035 Yuij Kohara unp
gb_est8: C08467	+	34.00	128.07	228.31	379	I C08467 C08467 Yuij Kohara unp

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LOCUS       D75467       335 bp      mRNA           EST           18-OCT-1999
DEFINITION  CELK104H2F Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
ACCESSION   D75467
VERSION     D75467.1 GI:1121251
SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE   1 (bases 1 to 335)
AUTHORS     Kohara,Y., Mitsuki,H., Nishigaki,A., Motohashi,T., Sugimoto,A. and
            Tabara,H.
TITLE       Toward an expression map of the C.elegans genome
JOURNAL     Unpublished (1994)
COMMENT     Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics
            Yata 1111, Mishima, Shizuoka 411, Japan
            Tel: 81-559-81-6854
            Fax: 81-559-81-6855
            Email: ykohara@lab.nig.ac.jp
            Insert Length: 651 Std Error: 0.00.
FEATURES             Location/Qualifiers
     source          1..335
                     /organism="Caenorhabditis elegans"
                     /strain="N2"
                     /db_xref="taxon:6239"
                     /clone="yk104h2"
                     /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
                     hermaphrodite embryo"
                     /sex="hermaphrodite"
                     /dev_stage="embryo"
BASE COUNT      94 a      84 c      74 g      80 t      3 others
ORIGIN
1 TyArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
260 TATTCCTTTCATTCGGATGATGAACGC 289

seq_name: gb_est5:D75467

alignment_scores:    Quality: 37.00      Length: 10
                    Ratio: 4.111      Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-13 x D75467

Align seg 1/1 to: D75467 from: 1 to: 335

1 TyArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
260 TATTCCTTTCATTCGGATGATGAACGC 289

seq_name: gb_est5:D73508

seq_documentation_block:    360 bp      mRNA           EST           18-OCT-1999
LOCUS       D73508
DEFINITION  CELK051H2F Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
ACCESSION   D73508
VERSION     D73508.1 GI:1119294
SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE   1 (bases 1 to 360)
AUTHORS     Kohara,Y., Mitsuki,H., Nishigaki,A., Motohashi,T., Sugimoto,A. and
            Tabara,H.
TITLE       Toward an expression map of the C.elegans genome
JOURNAL     Unpublished (1994)
COMMENT     Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics
            Yata 1111, Mishima, Shizuoka 411, Japan
            Tel: 81-559-81-6854
            Fax: 81-559-81-6855
            Email: ykohara@lab.nig.ac.jp
            Insert Length: 743 Std Error: 0.00
            High quality sequence stop: 257.
FEATURES             Location/Qualifiers
     source          1..360
                     /organism="Caenorhabditis elegans"
                     /strain="N2"
                     /db_xref="taxon:6239"
                     /clone="yk60g3"
                     /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
                     hermaphrodite embryo"
                     /sex="hermaphrodite"
                     /dev_stage="embryo"
BASE COUNT      103 a      87 c      84 g      80 t      6 others
ORIGIN
1 TyArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
241 TATTCCTTTCATTCGGATGATGAACGC 270

```

```

Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp
Insert Length: 2140 Std Error: 0.00
High quality sequence stop: 258.
FEATURES             Location/Qualifiers
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                     /organism="Caenorhabditis elegans"
                     /strain="N2"
                     /db_xref="taxon:6239"
                     /clone="yk51h2"
                     /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
                     hermaphrodite embryo"
                     /sex="hermaphrodite"
                     /dev_stage="embryo"
BASE COUNT      108 a      88 c      83 g      79 t      2 others
ORIGIN
1 TyArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
241 TATTCCTTTCATTCGGATGATGAACGC 270

seq_name: gb_est5:D73590

seq_documentation_block:    360 bp      mRNA           EST           18-OCT-1999
LOCUS       D73590
DEFINITION  CELK060G3F Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
ACCESSION   D73590
VERSION     D73590.1 GI:1119366
SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE   1 (bases 1 to 360)
AUTHORS     Kohara,Y., Mitsuki,H., Nishigaki,A., Motohashi,T., Sugimoto,A. and
            Tabara,H.
TITLE       Toward an expression map of the C.elegans genome
JOURNAL     Unpublished (1994)
COMMENT     Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics
            Yata 1111, Mishima, Shizuoka 411, Japan
            Tel: 81-559-81-6854
            Fax: 81-559-81-6855
            Email: ykohara@lab.nig.ac.jp
            Insert Length: 743 Std Error: 0.00
            High quality sequence stop: 257.
FEATURES             Location/Qualifiers
     source          1..360
                     /organism="Caenorhabditis elegans"
                     /strain="N2"
                     /db_xref="taxon:6239"
                     /clone="yk60g3"
                     /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
                     hermaphrodite embryo"
                     /sex="hermaphrodite"
                     /dev_stage="embryo"
BASE COUNT      103 a      87 c      84 g      80 t      6 others
ORIGIN
1 TyArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
241 TATTCCTTTCATTCGGATGATGAACGC 270

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ORIGIN

alignment_scores: Quality: 37.00 Length: 10
 Ratio: 4.111 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
 US-08-653-294-13 x D73590 ..

Align seg 1/1 to: D73590 from: 1 to: 360

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1 TytArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
258 TATTCCTTGTGATCGGATGGAACGC 287

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seq_name: gb_est5:D74728

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seq_documentation_block:
LOCUS D74728 360 bp mRNA EST 18-OCT-1999
DEFINITION CELK084F3F Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
embryo Caenorhabditis elegans cDNA clone yk84f3 5', mRNA sequence.
ACCESSION D74728
VERSION D74728
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE 1 (bases 1 to 360)
AUTHORS Kohara,Y., Mitsuki,H., Nishigaki,A., Motohashi,T., Sugimoto,A. and
Tabara,H.
TITLE Toward an expression map of the C.elegans genome
JOURNAL Unpublished (1994)
COMMENT On Sep 21, 1992 this sequence version replaced gi:276173.
Contact: Yuji Kohara
Gene Library Lab
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp
High quality sequence stop: 296.

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FEATURES

source
 1..360
 /organism="Caenorhabditis elegans"
 /strain="N2"
 /db_xref="taxon:6239"
 /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
 hermaphrodite embryo"
 /sex="hermaphrodite"
 /dev_stage="embryo"
 BASE COUNT 105 a 88 c 85 g 79 t 3 others
 ORIGIN

alignment_scores: Quality: 37.00 Length: 10
 Ratio: 4.111 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
 US-08-653-294-13 x D74728 ..

Align seg 1/1 to: D74728 from: 1 to: 360

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1 TytArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
257 TATTCCTTGTGATCGGATGGAACGC 286

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seq_name: gb_est5:D75423

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seq_documentation_block:
LOCUS D75423 360 bp mRNA EST 18-OCT-1999
DEFINITION CELK104C7F Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
embryo Caenorhabditis elegans cDNA clone yk104c7 5', mRNA sequence.
ACCESSION D75423
VERSION D75423
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE 1 (bases 1 to 360)
AUTHORS Kohara,Y., Mitsuki,H., Nishigaki,A., Motohashi,T., Sugimoto,A. and
Tabara,H.
TITLE Toward an expression map of the C.elegans genome
JOURNAL Unpublished (1994)
COMMENT Contact: Yuji Kohara
Gene Library Lab
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp
Insert Length: 1013 Std Error: 0.00
High quality sequence stop: 120.

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FEATURES

source
 1..360
 /organism="Caenorhabditis elegans"
 /strain="N2"
 /db_xref="taxon:6239"
 /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
 hermaphrodite embryo"
 /sex="hermaphrodite"
 /dev_stage="embryo"
 BASE COUNT 106 a 87 c 83 g 79 t 5 others
 ORIGIN

alignment_scores: Quality: 37.00 Length: 10
 Ratio: 4.111 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
 US-08-653-294-13 x D75423 ..

Align seg 1/1 to: D75423 from: 1 to: 360

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1 TytArgLeuAlaIleArgLeuAspGluArg 10
||| |||:|||||:|||||
248 TATTCCTTGTGATCGGATGGAACGC 277

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seq_name: gb_est5:D75727

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seq_documentation_block:
LOCUS D75727 360 bp mRNA EST 18-OCT-1999
DEFINITION CELK109D2F Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
embryo Caenorhabditis elegans cDNA clone yk109d2 5', mRNA sequence.
ACCESSION D75727
VERSION D75727
KEYWORDS EST.
SOURCE Caenorhabditis elegans.
ORGANISM Caenorhabditis elegans.
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE 1 (bases 1 to 360)
AUTHORS Kohara,Y., Mitsuki,H., Nishigaki,A., Motohashi,T., Sugimoto,A. and
Tabara,H.
TITLE Toward an expression map of the C.elegans genome
JOURNAL Unpublished (1994)
COMMENT Contact: Yuji Kohara

```

```

Gene Library Lab
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp
Insert Length: 1157 Std Error: 0.00.

FEATURES
    source
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            /organism="Caenorhabditis elegans"
            /strain="N2"
            /db_xref="taxon:6239"
            /clone_lib="Yui Kohara unpublished cDNA:Strain N2
            hermaphrodite embryo"
            /sex="hermaphrodite"
            /dev_stage="embryo"
BASE COUNT      105 a      87 c      75 t      8 others
ORIGIN

alignment_scores:
    Quality: 37.00      Length: 10
    Ratio: 4.111      Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
    US-08-653-294-13 x D75727      ..
    Align seg 1/1 to: D75727 from: 1 to: 360
        1 TyArgLeuAlaIleArgLeuAspGluArg 10
        ||| |||:|||||:|||||:|||||
        257 TATTCCTTCGATCGATGGAACGC 286

seq_name: gb_est16:C40575

seq_documentation_block:
    LOCUS      C40575      360 bp      mRNA      18-OCT-1999
    DEFINITION      C40575 Yui Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
    Caenorhabditis elegans cDNA clone yk246c11 5', mRNA sequence.
    ACCESSION      C40575
    VERSION      C40575.1 GI:2376812
    KEYWORDS      EST.
    SOURCE      Caenorhabditis elegans.
    ORGANISM      Caenorhabditis elegans.
    Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
    Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
    REFERENCE      1 (bases 1 to 360)
    AUTHORS      Kohara,Y., Motohashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
    Sano,M., Miyata,A. and Nishigaki,A.
    TITLE      Expression map of the C.elegans genome
    JOURNAL      Unpublished (1996)
    COMMENT      On Sep 12, 1996 this sequence version replaced gi:1395034.

FEATURES
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        1..360
            /organism="Caenorhabditis elegans"
            /strain="N2"
            /db_xref="taxon:6239"
            /clone_lib="Yui Kohara unpublished cDNA:Strain N2
            hermaphrodite embryo"
            /sex="hermaphrodite"
            /dev_stage="embryo"
BASE COUNT      109 a      93 c      75 t      1 others
ORIGIN

alignment_scores:
    Quality: 37.00      Length: 10
    Ratio: 4.111      Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
    US-08-653-294-13 x C42212      ..
    Align seg 1/1 to: C42212 from: 1 to: 360
        1 TyArgLeuAlaIleArgLeuAspGluArg 10
        ||| |||:|||||:|||||:|||||
        222 TATTCCTTCGATCGATGGAACGC 251

seq_name: gb_est16:C44981

seq_documentation_block:
    LOCUS      C40575      360 bp      mRNA      18-OCT-1999
    DEFINITION      C40575 Yui Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
    Caenorhabditis elegans cDNA clone yk246c11 5', mRNA sequence.
    ACCESSION      C40575
    VERSION      C40575.1 GI:2376812
    KEYWORDS      EST.
    SOURCE      Caenorhabditis elegans.
    ORGANISM      Caenorhabditis elegans.
    Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
    Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
    REFERENCE      1 (bases 1 to 360)
    AUTHORS      Kohara,Y., Motohashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
    Sano,M., Miyata,A. and Nishigaki,A.
    TITLE      Expression map of the C.elegans genome
    JOURNAL      Unpublished (1996)
    COMMENT      On Sep 12, 1996 this sequence version replaced gi:1395034.

FEATURES
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            /db_xref="taxon:6239"
            /clone_lib="Yui Kohara unpublished cDNA:Strain N2
            hermaphrodite embryo"
            /sex="hermaphrodite"
            /dev_stage="embryo"
BASE COUNT      109 a      82 g      78 t      1 others
ORIGIN

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alignment_scores:
    Quality: 37.00      Length: 10
    Ratio: 4.111      Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
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        1 TyArgLeuAlaIleArgLeuAspGluArg 10
        ||| |||:|||||:|||||:|||||
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seq_name: gb_est16:C42212

seq_documentation_block:
    LOCUS      C42212      360 bp      mRNA      18-OCT-1999
    DEFINITION      C42212 Yui Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
    Caenorhabditis elegans cDNA clone yk290d10 5', mRNA sequence.
    ACCESSION      C42212
    VERSION      C42212.1 GI:2378449
    KEYWORDS      EST.
    SOURCE      Caenorhabditis elegans.
    ORGANISM      Caenorhabditis elegans.
    Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
    Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
    REFERENCE      1 (bases 1 to 360)
    AUTHORS      Kohara,Y., Motohashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
    Sano,M., Miyata,A. and Nishigaki,A.
    TITLE      Expression map of the C.elegans genome
    JOURNAL      Unpublished (1996)
    COMMENT      On Sep 12, 1996 this sequence version replaced gi:1400867.

FEATURES
    source
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            /strain="N2"
            /db_xref="taxon:6239"
            /clone_lib="Yui Kohara unpublished cDNA:Strain N2
            hermaphrodite embryo"
            /sex="hermaphrodite"
            /dev_stage="embryo"
BASE COUNT      109 a      90 c      82 g      78 t      1 others
ORIGIN

alignment_scores:
    Quality: 37.00      Length: 10
    Ratio: 4.111      Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
    US-08-653-294-13 x C42212      ..
    Align seg 1/1 to: C42212 from: 1 to: 360
        1 TyArgLeuAlaIleArgLeuAspGluArg 10
        ||| |||:|||||:|||||:|||||
        222 TATTCCTTCGATCGATGGAACGC 251

seq_name: gb_est16:C44981

seq_documentation_block:
    LOCUS      C42212      360 bp      mRNA      18-OCT-1999
    DEFINITION      C42212 Yui Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
    Caenorhabditis elegans cDNA clone yk290d10 5', mRNA sequence.
    ACCESSION      C42212
    VERSION      C42212.1 GI:2378449
    KEYWORDS      EST.
    SOURCE      Caenorhabditis elegans.
    ORGANISM      Caenorhabditis elegans.
    Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
    Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
    REFERENCE      1 (bases 1 to 360)
    AUTHORS      Kohara,Y., Motohashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
    Sano,M., Miyata,A. and Nishigaki,A.
    TITLE      Expression map of the C.elegans genome
    JOURNAL      Unpublished (1996)
    COMMENT      On Sep 12, 1996 this sequence version replaced gi:1400867.

FEATURES
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            /db_xref="taxon:6239"
            /clone_lib="Yui Kohara unpublished cDNA:Strain N2
            hermaphrodite embryo"
            /sex="hermaphrodite"
            /dev_stage="embryo"
BASE COUNT      109 a      90 c      82 g      78 t      1 others
ORIGIN

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LOCUS       C44981               360 bp      mRNA      EST      18-OCT-1999
DEFINITION  C44981 Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
            Caenorhabditis elegans cDNA clone yk373f5 5', mRNA sequence.
ACCESSION   C44981
VERSION     C44981.1
KEYWORDS    C44981.1 GI:2381218
SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Caenorhabditis elegans
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE   1 (bases 1 to 360)
AUTHORS     Kohara,Y., Mochashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
            Sano,M., Miyata,A. and Nishigaki,A.
TITLE       Expression map of the C.elegans genome
JOURNAL     Unpublished (1996)
COMMENT     On May 8, 1995 this sequence version replaced gi:801522.
            Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics
            Yata 1111, Mishima, Shizuoka 411, Japan
            Tel: 81-559-81-6854
            Fax: 81-559-81-6855
            Email: ykohara@lab.nig.ac.jp.

FEATURES             Location/Qualifiers
     source           1..360
                     /organism="Caenorhabditis elegans"
                     /strain="N2"
                     /db_xref="taxon:6239"
                     /clone="yk373f5"
                     /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
                     hermaphrodite embryo"
                     /sex="hermaphrodite"
                     /dev_stage="embryo"

BASE COUNT      107 a      95 c      79 g      75 t      4 others
ORIGIN
1  TyArgLeuAlaIleArgLeuAspGluArg 10
|||||
210 TATTCTTTTCATTCGATCGATGAACGC 239

alignment_scores:
  Quality: 37.00      Length: 10
  Ratio: 4.111      Gaps: 0
  Percent Similarity: 90.000      Percent Identity: 70.000

alignment_block:
US-08-653-294-13 x C44981 ..
Align seg 1/1 to: C44981 from: 1 to: 360

seq_name: gb_est16:C51139

seq_documentation_block:
LOCUS       C51139               360 bp      mRNA      EST      18-OCT-1999
DEFINITION  C51139 Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
            Caenorhabditis elegans cDNA clone yk491f7 5', mRNA sequence.
ACCESSION   C51139
VERSION     C51139.1
KEYWORDS    C51139.1 GI:2388392
SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Caenorhabditis elegans
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
            Kohara,Y., Mochashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
            Sano,M., Miyata,A. and Nishigaki,A.
            Expression map of the C.elegans genome
            Unpublished (1996)
            On Sep 12, 1996 this sequence version replaced gi:1395322.
            Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics

seq_name: gb_est16:C51139

seq_documentation_block:
LOCUS       C51139               360 bp      mRNA      EST      18-OCT-1999
DEFINITION  C51139 Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite embryo
            Caenorhabditis elegans cDNA clone yk491f7 5', mRNA sequence.
ACCESSION   C51139
VERSION     C51139.1
KEYWORDS    C51139.1 GI:2388392
SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Caenorhabditis elegans
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
            Kohara,Y., Mochashi,T., Tabara,H., Watanabe,H., Sugimoto,A.,
            Sano,M., Miyata,A. and Nishigaki,A.
            Expression map of the C.elegans genome
            Unpublished (1996)
            On Sep 12, 1996 this sequence version replaced gi:1395322.
            Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics

```

```

Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.

FEATURES             Location/Qualifiers
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            embryo Caenorhabditis elegans cDNA clone yk607d3 5', mRNA sequence.
ACCESSION   AV192603
VERSION     AV192603.1
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SOURCE      EST.
ORGANISM    Caenorhabditis elegans.
            Caenorhabditis elegans
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
            Kohara,Y., Shin-I,T., Thierry-Mieg,J., Thierry-Mieg,D., Mitsuki,H.,
            Nishigaki,A., Mochashi,T., Zeng,Q., Watanabe,H., Sugimoto,A.,
            Sano,M., Miyata,A., Mitani,Y., Iida,K., Uesugi,H., Sugiyama,Y. and
            Nomoto,H.
            Expressed genes in C.elegans
            Unpublished (1999)
            On Jun 22, 1998 this sequence version replaced gi:3247424.
            Contact: Yuji Kohara
            Gene Library Lab
            National Institute of Genetics
            Yata 1111, Mishima, Shizuoka 411, Japan
            Tel: 81-559-81-6854
            Fax: 81-559-81-6855
            Email: ykohara@lab.nig.ac.jp.

FEATURES             Location/Qualifiers
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  embryo Caenorhabditis elegans cDNA clone yk609e10 5', mRNA
  sequence.
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  ORGANISM Caenorhabditis elegans.
  Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
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  Kohara,Y., Shin-i,T., Thierry-Mieg,J., Thierry-Mieg,D., Mitsuiki,H.,
  Nishigaki,A., Motohashi,T., Zeng,Q., Watanabe,H., Sugimoto,A.,
  Sano,M., Miyata,A., Mitani,Y., Iida,K., Uesugi,H., Sugiyama,Y. and
  Nomoto,H.
  Expressed genes in C.elegans
  Unpublished (1999)
  On Feb 18, 1999 this sequence version replaced gi:4299301.
  Contact: Yuji Kohara
  Gene Library Lab
  National Institute of Genetics
  Yata 1111, Mishima, Shizuoka 411, Japan
  Tel: 81-559-81-6854
  Fax: 81-559-81-6855
  Email: ykohara@lab.nig.ac.jp.
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  sequence.
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  VERSION AV193922
  KEYWORDS EST.
  SOURCE Caenorhabditis elegans.
  ORGANISM Caenorhabditis elegans.
  Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
  Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
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  Kohara,Y., Shin-i,T., Thierry-Mieg,J., Thierry-Mieg,D., Mitsuiki,H.,
  Nishigaki,A., Motohashi,T., Zeng,Q., Watanabe,H., Sugimoto,A.,
  Sano,M., Miyata,A., Mitani,Y., Iida,K., Uesugi,H., Sugiyama,Y. and
  Nomoto,H.
  Expressed genes in C.elegans
  Unpublished (1999)
  On Mar 10, 1998 this sequence version replaced gi:2948755.
  Contact: Yuji Kohara
  Gene Library Lab
  National Institute of Genetics
  Yata 1111, Mishima, Shizuoka 411, Japan
  Tel: 81-559-81-6854
  Fax: 81-559-81-6855
  Email: ykohara@lab.nig.ac.jp.
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  DEFINITION AV194609 Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
  embryo Caenorhabditis elegans cDNA clone yk631f9 5', mRNA
  sequence.
  ACCESSION AV194609
  VERSION AV194609.1 GI:5576761
  KEYWORDS EST.
  SOURCE Caenorhabditis elegans.
  ORGANISM Caenorhabditis elegans.
  Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
  Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
  1 (bases 1 to 360)
  Kohara,Y., Shin-i,T., Thierry-Mieg,J., Thierry-Mieg,D., Mitsuiki,H.,
  Nishigaki,A., Motohashi,T., Zeng,Q., Watanabe,H., Sugimoto,A.,
  Sano,M., Miyata,A., Mitani,Y., Iida,K., Uesugi,H., Sugiyama,Y. and
  Nomoto,H.

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TITLE Expressed genes in C.elegans
 JOURNAL Unpublished (1999)
 COMMENT On May 9, 1996 this sequence version replaced gi:1132813.
 Contact: Yuji Kohara
 Gene Library Lab
 National Institute of Genetics
 Yata 1111, Mishima, Shizuoka 411, Japan
 Tel: 81-559-81-6854
 Fax: 81-559-81-6855
 Email: ykohara@lab.nig.ac.jp.
 Location/Qualifiers
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 /clone="yk631f9"
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 239 TATTCTCTTCGATCGATCGATGACGC 268

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2000, 01:29:38 ; Search time 122.56 Seconds
(without alignments)
1.933 Million cell updates/sec

Title: US-08-653-294-14
Perfect score: 49
Sequence: 1 REDRLILLY 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : A_Geneseq_36.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	49	100.0	10	1 W47271	Immunomodulatory p
2	44	89.8	10	1 R41212	Peptide fragment o
3	44	89.8	10	1 R83075	HLA-B2702 CTL modu
4	44	89.8	10	1 R83094	HLA-B2702 CTL modu
5	44	89.8	10	1 R83096	HLA-B2702 CTL modu
6	44	89.8	10	1 R95423	HLA-B2705.75-84. C
7	44	89.8	10	1 R95425	HLA-B2702.75-84(D)
8	44	89.8	10	1 W07513	T-cell modulating
9	44	89.8	10	1 W47267	Immunomodulatory p
10	44	89.8	10	1 W47269	Immunomodulatory p
11	44	89.8	10	1 W33785	Peptide B2705.75-8
12	44	89.8	10	1 W33787	Peptide B2702.75-8
13	44	89.8	10	1 W33789	Peptide B2702.75-8
14	44	89.8	17	1 R71442	Human HLA-B27-(62-
15	44	89.8	17	1 R71443	Human [Phe4]-HLA-
16	44	89.8	25	1 R41221	Peptide fragment o
17	44	89.8	25	1 R83091	HLA-B2702 CTL modu
18	44	89.8	25	1 R95417	HLA-B2705.60-84. C
19	44	89.8	337	1 P70590	Sequence of the hu
20	44	89.8	362	1 P70155	Sequence encoded b
21	39	79.6	10	1 R41208	Peptide fragment o
22	39	79.6	10	1 R83062	HLA-B2702 CTL modu
23	39	79.6	10	1 R95413	Alpha1-helix of HL
24	39	79.6	10	1 R95427	HLA-B2702.75-84(L)
25	39	79.6	10	1 W07512	T-cell modulating
26	39	79.6	10	1 W07514	T-cell modulating
27	39	79.6	10	1 W47265	Immunomodulatory p
28	39	79.6	10	1 W33784	Peptide B2702.75-8
29	39	79.6	15	1 R92912	HLA-B2702 CTL modu
30	39	79.6	15	1 W33795	Peptide B2702.70-8
31	39	79.6	20	1 R92907	HLA-B2702 CTL modu
32	39	79.6	20	1 R92908	HLA-B2702 CTL modu
33	39	79.6	20	1 R95428	HLA-B2702 84-75-84
34	39	79.6	20	1 W33778	Immunomodulating d

35 39 79.6 20 1 W33791 Peptide B2702.84-7
36 39 79.6 25 1 R41205 Peptide fragment o
37 39 79.6 25 1 R48286 Peptide fragment o
38 39 79.6 25 1 R83090 HLA-B2702 CTL modu
39 39 79.6 25 1 R83093 HLAB38 CTL modulac
40 39 79.6 25 1 R95416 HLA-B2702.60-84. C
41 39 79.6 25 1 R95422 HLAB38.6084. Comps
42 39 79.6 25 1 W33794 Peptide B2702.60-8
43 39 79.6 184 1 Y06801 Peptide Seq ID No:
44 39 79.6 362 1 R03142 Sequence of HLA-B*
45 39 79.6 362 1 R03144 Sequence of HLA-B5

ALIGNMENTS

RESULT 1

W47271
ID W47271 standard; peptide; 10 AA.
AC W47271;
DT 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.
FH Key Location/Qualifiers
FT Misc_difference 1. .10 /note= "at least one of the amino acids is the D-isomer"
FT

PN W09744052-A1.
PD 27-NOV-1997.
PF 23-APR-1997; U06705.
PR 22-MAY-1996; US-651650.
PA (STRD) UNIV IELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 98-018220/02.

PT Novel immunomodulatory peptide-type compound - useful for inhibiting transplant rejection
PS Claim 10; Page 36; 41pp; English.
CC The present sequence is an immunomodulatory peptide, which comprises a Class I HLA-B alpha-1 domain sequence. It can be used in a pharmaceutical composition together with a subtherapeutic dose of an immunosuppressant, to extend the period of acceptance of a transplant from a major histocompatibility complex (MHC) unmatched donor, i.e. to inhibit transplant rejection. It can also be used in the treatment of autoimmune diseases.
CC Peptides using the D-form amino acids are more effective immunomodulators than their diastereomers or enantiomers.
SQ Sequence 10 AA;

Query Match 100.0%; Score 49; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 0.0009;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 REDRLILLY 10
| | | | | | | | | |
Db 1 REDRLILLY 10

RESULT 2

R41212
ID R41212 standard; peptide; 10 AA.
AC R41212;
DT 15-MAR-1994 (first entry)
DE Peptide fragment of Class I HLA peptide.
KW Human leukocyte antigen; HLA; peptide; transplantation; neoplasia;
KW parasitic disease; cytotoxic T lymphocyte; modulation.
OS Synthetic.
PN W09317699-A.
PD 16-SEP-1993.
PF 25-FEB-1993; U01758.

PR 02-MAR-1992; US-844716.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger CA, Krensky AM;
 DR WPI: 93-303134/38.
 PT New peptide(s) Based on Class I HLA antigen domains - used for
 PS modulating cytotoxic T-lymphocyte activity towards targets
 PT Claim 11; Page 54; 61pp; English.
 CC The peptide is used to modulate cytotoxic T-lymphocyte (CTL)
 CC activity, either by inhibition or stimulation. It can be used
 CC for inhibiting CTL toxicity in transplantations, for inducing CTL
 CC activity in parasitic diseases and neoplasia and in studies on viral
 CC infection. The peptide can also be used for identifying CTLs which
 CC bind to it and removing subsets of CTLs from a T-cell composition.
 CC This peptide sequence is more commonly found within larger peptide
 CC compounds of not more than 30 amino acids in length.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLRY 10
 DB 1 REDRLRLRY 10

RESULT 3
 R83075
 ID R83075 standard; peptide; 10 AA.
 AC R83075;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.75-84).
 CC Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN W09526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Farham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host
 PS Example 14; Page 34; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC corresponds to residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLRY 10
 DB 1 REDRLRLRY 10

RESULT 4
 R83094

ID R83094 standard; peptide; 10 AA.
 AC R83094;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.75-84(D)).
 CC Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN W09526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Farham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host
 PS Example 14; Page 34; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC corresponds to residues 75-84 of the alpha-1 domain of the class I MHC
 CC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with
 CC a subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLRY 10
 DB 1 REDRLRLRY 10

RESULT 5
 R83096
 ID R83096 standard; peptide; 10 AA.
 AC R83096;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.75-84(L)).
 CC Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN W09526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Farham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host
 PS Example 14; Page 34; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC corresponds to residues 75-84 of the alpha-1 domain of the class I MHC
 CC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with
 CC a subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 10 AA;

CC of the patient.
SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.0084;
Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
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Db 1 RENLRLRLRY 10

RESULT 6

R95423 ID R95423 standard; peptide; 10 AA.
AC R95423;
DT 12-NOV-1996 (first entry)
DE HLA-B*2705:75-84.
KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN W09513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C. Krensky AM;
DR WPI; 95-194027/25.
PT Compsns. comprising lymphoid surface membrane proteins - which may
PT inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 11; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
CC human-leucocyte-associated antigens. This sequence represents the
CC HLA-B*2705:75-84. These sequences can be used to isolate the protein
CC from a T-cell lysate. p74 is a T-cell surface membrane protein
CC associated with T-cell activation in mammalian T-cells, and is also
CC immunologically cross reactive with the heat shock protein Hsc70. p74 is
CC found in a limited number of cell types, but is particularly expressed on
CC B and T cells. p74 can be isolated by lysis of a suitable cell with an
CC amphoteric detergent, and then passed through an affinity column
CC containing a covalently bound HLA-B*2702 palindromic peptide.
CC Compositions comprising the extracellular fragment of p74 combined with
CC HLA-B*2702:60-84 (see R95416), induces calcium influx, and inhibits
CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
CC compounds can be screened for their effect on the cytolytic activity of
CC T-cells, by combining them with the extracellular portion of p74 and
CC determining the amount of binding between the candidate compound and p74.
CC Modulation of CTL activity can be inhibited in a cellular composition
CC containing T-cells and antigen presenting cells (APCs), by adding to the
CC mix the extracellular portion of p74, in an amount sufficient to compete
CC with p74 for the binding of the p74 ligand.
SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.0084;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
|||||
Db 1 REDRLRLRY 10

RESULT 7

R95425 ID R95425 standard; peptide; 10 AA.
AC R95425;
DT 12-NOV-1996 (first entry)
DE HLA-B*2702:75-84(D).
KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;

KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
OS Synthetic.
EH Key Location/Qualifiers
FT misc_difference 3 /note= "N3D mutation"
PN W09513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C. Krensky AM;
DR WPI; 95-194027/25.
PT Compsns. comprising lymphoid surface membrane proteins - which may
PT inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 11; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
CC human-leucocyte-associated antigens. This sequence represents the
CC HLA-B*2702:75-84(D). These sequences can be used to isolate the protein
CC from a T-cell lysate. p74 is a T-cell surface membrane protein
CC associated with T-cell activation in mammalian T-cells, and is also
CC immunologically cross reactive with the heat shock protein Hsc70. p74 is
CC found in a limited number of cell types, but is particularly expressed on
CC B and T cells. p74 can be isolated by lysis of a suitable cell with an
CC amphoteric detergent, and then passed through an affinity column
CC containing a covalently bound HLA-B*2702 palindromic peptide.
CC Compositions comprising the extracellular fragment of p74 combined with
CC HLA-B*2702:60-84 (see R95416), induces calcium influx, and inhibits
CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
CC compounds can be screened for their effect on the cytolytic activity of
CC T-cells, by combining them with the extracellular portion of p74 and
CC determining the amount of binding between the candidate compound and p74.
CC Modulation of CTL activity can be inhibited in a cellular composition
CC containing T-cells and antigen presenting cells (APCs), by adding to the
CC mix the extracellular portion of p74, in an amount sufficient to compete
CC with p74 for the binding of the p74 ligand.
SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.0084;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
|||||
Db 1 REDRLRLRY 10

RESULT 8

W07513 ID W07513 standard; peptide; 10 AA.
AC W07513;
DT 04-AUG-1997 (first entry)
DE T-cell modulating peptide #2.
KW T-cell modulator; autoimmune disease; tissue destruction; alpha1-domain;
KW mammal; major histocompatibility complex; MHC class I; antigen; perforin;
KW insulin-dependent diabetes mellitus; multiple sclerosis; inflammation;
KW rheumatoid arthritis; psoriasis; pemphigus vulgaris; Sjogren's disease;
KW thyroid disease; Hashimoto's thyroiditis; myasthenia gravis; granzyme;
KW autologous target cell; cytokine release; T cell activation; therapy.
OS Synthetic.
PN W09635443-A1.
PD 14-NOV-1996.
PF 05-APR-1996; U04710.
PR 12-MAY-1995; US-440504.
PA (SANG-) SANGSTAT MEDICAL CORP.
PI Buelow R;
DR WPI; 96-518410/51.
PT Treatment of auto-immune disease by admin. of peptide(s) corresp. to
PT major histocompatibility complex antigens - esp. for delaying onset
PT of clinical symptoms of insulin dependent diabetes by modulating T
PT cell mediated attack on target cells

PS Claim 7; Page 20; 24pp; English.
 CC W07512-W07518 represent T-cell modulating peptides that can be used in
 CC the method of the invention. These sequences are based on a portion of
 CC the generic peptide corresponding to residues 70-91 of the alpha-1 domain
 CC of the major histocompatibility complex (MHC) class I antigen (see
 CC W07510). The method is for affecting the course of an autoimmune disease
 CC involving T-cell mediated destruction of tissue in mammals. These
 CC peptides are used especially to treat insulin-dependent diabetes
 CC mellitus, preferably being administered during the pre-clinical stage to
 CC delay onset of the disease. Other diseases that can be treated are
 CC multiple sclerosis, rheumatoid arthritis, psoriasis, pemphigus vulgaris,
 CC Sjogren's disease, thyroid disease, Hashimoto's thyroiditis, myasthenia
 CC gravis, etc. The peptides modulate T-cell mediated attack on autologous
 CC target cells, and may also reduce inflammation, swelling, and release of
 CC cytokines, perforins, granzymes etc. associated with T cell activation.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLILRY 10
 | | | | |
 Db 1 REDRLIALRY 10

RESULT 9

W47267 ID W47267 standard; peptide; 10 AA.
 AC W47267;
 DT 22-MAY-1998 (first entry)
 DE Immunomodulatory peptide.
 KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
 OS Homo sapiens.
 OS Synthetic.
 FH Key Location/Qualifiers
 FT Misc_difference 1..10 /note= "at least one of the amino acids is the
 FT D-isomer

PN W09744052-A1.
 PD 27-NOV-1997.
 PF 23-APR-1997; U06705.
 PR 22-MAY-1996; US-651650.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI; 98-018220/02.
 PT Novel immunomodulatory peptide-type compound - useful for inhibiting
 PT transplant rejection
 PS Claim 10; Page 36; 41pp; English.
 CC The present sequence is an immunomodulatory peptide, which
 CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
 CC in a pharmaceutical composition together with a subtherapeutic dose
 CC of an immunosuppressant, to extend the period of acceptance of a
 CC transplant from a major histocompatibility complex (MHC) unmatched
 CC donor, i.e. to inhibit transplant rejection. It can also be used in
 CC the treatment of autoimmune diseases.
 CC Peptides using the D-form amino acids are more effective
 CC immunomodulators than their diastereomers or enantiomers.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 REDRLILRY 10
 | | | | |
 Db 1 REDRLIALRY 10

RESULT 10

W47269 ID W47269 standard; peptide; 10 AA.
 AC W47269;
 DT 22-MAY-1998 (first entry)
 DE Immunomodulatory peptide.
 KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
 OS Homo sapiens.
 OS Synthetic.
 FH Key Location/Qualifiers
 FT Misc_difference 1..10 /note= "at least one of the amino acids is the
 FT D-isomer
 PN W09744052-A1.
 PD 27-NOV-1997.
 PF 23-APR-1997; U06705.
 PR 22-MAY-1996; US-651650.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI; 98-018220/02.
 PT Novel immunomodulatory peptide-type compound - useful for inhibiting
 PT transplant rejection
 PS Claim 10; Page 36; 41pp; English.
 CC The present sequence is an immunomodulatory peptide, which
 CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
 CC in a pharmaceutical composition together with a subtherapeutic dose
 CC of an immunosuppressant, to extend the period of acceptance of a
 CC transplant from a major histocompatibility complex (MHC) unmatched
 CC donor, i.e. to inhibit transplant rejection. It can also be used in
 CC the treatment of autoimmune diseases.
 CC Peptides using the D-form amino acids are more effective
 CC immunomodulators than their diastereomers or enantiomers.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLILRY 10
 | | | | |
 Db 1 REDRLIALRY 10

RESULT 11
 W33785 ID W33785 standard; peptide; 10 AA.
 AC W33785;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2705.75-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI; 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or

CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDLRLLRY 10
 ||||| ||||
 Db 1 REDLRLLRY 10

RESULT 12

W33787
 ID W33787 standard; peptide; 10 AA.
 AC W33787;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.75-84D77 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W05744351-A1.
 PD 27-NOV-1997.
 PF 27-NOV-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.

PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDLRLLRY 10
 ||||| ||||
 Db 1 REDLRLLRY 10

RESULT 12
 W33787
 ID W33787 standard; peptide; 10 AA.
 AC W33787;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.75-84D77 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W05744351-A1.
 PD 27-NOV-1997.
 PF 27-NOV-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.

PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 10 AA;

Db 1 REDLRLLRY 10
 ||||| ||||

RESULT 13

W33789
 ID W33789 standard; peptide; 10 AA.
 AC W33789;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.75-84L81 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W05744351-A1.
 PD 27-NOV-1997.
 PF 27-NOV-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.

PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 10 AA;

Query Match 89.8%; Score 44; DB 1; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.0084;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 1 REDLRLLRY 10
 ||||| ||||
 Db 1 REDLRLLRY 10

RESULT 14

R71442
 ID R71442 standard; peptide; 17 AA.
 AC R71442;
 DT 12-OCT-1995 (first entry)
 DE Human HLA-B27-(62-85) antigen derived peptide.
 KW Human HLA-B27-(62-85) antigen derived peptide;
 KW interaction modulation; arthritis; neoplasias; lupus erythematosus.
 OS Homo sapiens.
 PN W0505189-A.
 PD 23-FEB-1995.
 PF 12-AUG-1994; U09189.
 PR 12-AUG-1993; US-105416.
 PA (REGC) UNIV CALIFORNIA.
 PI Goldstein A, Goodenow RS, Olsson L;
 DR WPI: 95-098577/13.

PT Regulating cell surface receptor response - by modulating
 PT interaction between MHC class I antigen and the cell surface
 PT receptor
 PS Example 4; Page 45; 103pp; English.
 CC R71439-R71443 are human major histocompatibility complex class 1
 CC (MHC 1) alpha 1 domain and HLA derived peptides and fusion peptides.
 CC They were used to modulate interactions between MHC 1/HLA and cell
 CC surface receptors. Via competitive inhibition the peptides diminish
 CC the receptors response, this feature may be useful for the treatment
 CC of neoplasias, lupus erythematosus and arthritis.
 SQ Sequence 17 AA;

Query Match 89.8%; Score 44; DB 1; Length 17;
 Best Local Similarity 90.0%; Pred. No. 0.015;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
 ||||| ||||
 Db 7 REDLRILLRY 16

RESULT 15

R71443
 ID R71443 standard; peptide: 17 AA.
 AC R71443;
 DT 12-OCT-1995 (first entry)
 DE Human [Phe74]-HLA-B27-(62-85) antigen derived peptide.
 KW Human [Phe74]-HLA-B27-(62-85) antigen derived peptide; cell receptor;
 KW interaction modulation; arthritis; neoplasias; lupus erythematosus.
 OS Homo sapiens.
 PN W09505189-A.
 PD 23-FEB-1995.
 PF 12-AUG-1994; U09189.
 PR 12-AUG-1993; US-105416.
 PA (REGC) UNIV CALIFORNIA.
 PI Goldstein A, Goodenow RS, Olsson L;
 DR WPI; 95-098577/13.
 PT Regulating cell surface receptor response - by modulating
 PT interaction between MHC class I antigen and the cell surface
 PT receptor
 PS Example 4; Page 45; 103pp; English.
 CC R71439-R71443 are human major histocompatibility complex class 1
 CC (MHC 1) alpha 1 domain and HLA derived peptides and fusion peptides.
 CC They were used to modulate interactions between MHC 1/HLA and cell
 CC surface receptors. Via competitive inhibition the peptides diminish
 CC the receptors response, this feature may be useful for the treatment
 CC of neoplasias, lupus erythematosus and arthritis.
 SQ Sequence 17 AA;

Query Match 89.8%; Score 44; DB 1; Length 17;
 Best Local Similarity 90.0%; Pred. No. 0.015;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
 ||||| ||||
 Db 7 REDLRILLRY 16

Search completed: February 8, 2000, 01:29:38
 Job time: 1750 sec

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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:25 : Search time 117.7 seconds
(without alignments)
4.008 Million cell updates/sec

Title: US-08-653-294-14
Perfect score: 49
Sequence: 1 REDRLRLRY 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : PIR-62:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	44	89.8	338	2 I56116	MHC HLA-B27-HS - h
2	44	89.8	362	1 HLUHUB2	MHC class I histoc
3	44	89.8	362	2 C35997	MHC class I histoc
4	44	89.8	362	2 I68724	MHC class I histoc
5	44	89.8	362	2 I37485	human lymphocyte a
6	44	89.8	362	2 I54289	MHC HLA-B27d - hum
7	39	79.6	137	2 I80174	class I histocompa
8	39	79.6	273	2 I38509	MHC class I histoc
9	39	79.6	274	2 I54463	MHC HLA-B38 chain
10	39	79.6	334	2 I59308	class I histocompa
11	39	79.6	354	2 I80168	class I histocompa
12	39	79.6	354	2 I80167	class I histocompa
13	39	79.6	355	2 I80169	class I histocompa
14	39	79.6	355	2 I80171	class I histocompa
15	39	79.6	359	1 HLUH12	MHC class I histoc
16	39	79.6	362	1 HLUHUB8	MHC class I histoc
17	39	79.6	362	2 B30345	MHC class I histoc
18	39	79.6	362	2 JH0541	class I histocompa
19	39	79.6	362	2 JH0539	class I histocompa
20	39	79.6	362	2 JH0540	class I histocompa
21	39	79.6	362	2 A45834	MHC class I histoc
22	39	79.6	362	2 I84486	transmembrane glyc
23	39	79.6	362	2 I62045	gene HLA B-1517 pr
24	39	79.6	362	2 I84490	lymphocyte antigen
25	39	79.6	362	2 I37521	HLA-Bw57.2 antigen
26	39	79.6	362	2 A30345	MHC class I histoc
27	39	79.6	362	2 I59633	MHC HLA-B transmem
28	39	79.6	362	2 S24434	class I histocompa
29	39	79.6	362	2 I37120	MHC class I histoc
30	39	79.6	363	2 S07113	class I histocompa

31 39 79.6 363 2 S03537 class I histocompa
32 39 79.6 364 2 D35997 MHC class I histoc
33 39 79.6 365 2 S77963 MHC class I histoc
34 39 79.6 365 2 I54416 HLA-AW24 protein -
35 39 79.6 365 2 I54493 MHC class I histoc
36 38 77.6 362 1 HLUH32 MHC class I histoc
37 38 77.6 362 2 I37515 MHC class I histoc
38 38 77.6 364 2 A35997 MHC class I histoc
39 37 75.5 328 2 I54414 MHC H-2K transplan
40 37 75.5 362 2 I71998 MHC H-2D-k protein
41 37 75.5 368 2 I68705 MHC H-2K-w28 prote
42 34 69.4 339 2 T15113 hypothetical prote
43 34 69.4 355 2 I37516 HLA-B alpha-chain
44 34 69.4 358 1 ADMU fructose-bisphosph
45 34 69.4 358 2 T05052 fructose-bisphosph

ALIGNMENTS

RESULT 1

I56116
MHC HLA-B27-HS - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 23-Jul-1999
C:Accession: I56116
R:Choo, S.Y.; Fan, L.A.; Hansen, J.A.
J. Immunol. 147, 174-180, 1991
A:title: A novel HLA-B27 allele maps B27 allospecificity to the region around positio
A:Reference number: I56116; MUID:91268545
A:Accession: I56116
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-338 <RES>
A:Cross-references: GB:M62852; NID:gl87760; PIDN:AAA59647.1; PID:gl87761
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 89.8%; Score 44; DB 2; Length 338;
Best Local Similarity 90.0%; Pred. No. 0.2;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLRY 10

Db 75 REDRLRLRY 84

RESULT 2

HLHUB2
MHC class I histocompatibility antigen HLA-B27 alpha chain precursor - human
C:Species: Homo sapiens (man)
C:Date: 13-Apr-1986 #sequence_revision 28-Apr-1995 #text_change 22-Jun-1999
C:Accession: S07441; A25092; B25092; A94087; S34180; S44942; A90493; B24741; I55965;
R:Weiss, E.H.; Kuon, W.; Doerner, C.; Lang, M.; Riethmuller, G.
Immunobiology 170, 367-380, 1985
A:title: Organization, sequence and expression of the HLA-B27 gene: a molecular appro
A:Reference number: S07441; MUID:86138405
A:Accession: S07441
A:Molecule type: DNA
A:Residues: 1-362 <WEI>
A:Cross-references: EMBL:X03945
A:Note: the authors translated the codon GAC for residue 61 as Ala and the codon CAG
A:Note: this allele is designated B*27052 (formerly 27W)
R:Seemann, G.H.A.; Rein, R.S.; Brown, C.S.; Ploegh, H.L.
EMBO J. 5, 547-552, 1986
A:title: Gene conversion-like mechanisms may generate polymorphism in human class I g
A:Reference number: A91061; MUID:86220133
A:Accession: A25092
A:Molecule type: DNA
A:Residues: 1-362 <SEE>
A:Cross-references: GB:X03665; NID:g32250; PIDN:CAA27302.1; PID:g871297.
A:Note: this allele is designated B*27051 (formerly 27W)
A:Accession: B25092

A:Molecule type: DNA
A:Residues: 1-100,'N','102-103','IA',106-362 <SE2>
A:Cross-references: GB:X03664; NID:g32236; PIDN:CAA27301.1; PID:g871296
A:Note: this allele is designated B*2702 (formerly 27K)
R:Znots, H.; Rietmueller, G.; Weiss, E.; Meo, F.
Proc. Natl. Acad. Sci. U.S.A. 83, 1428-1432, 1986
A:Title: Complete sequence of HLA-B*27 cDNA identified through the characterization of s
A:Reference number: A34087; MUID:86149317
A:Accession: A94087
A:Molecule type: mRNA
A:Residues: 25-205,'V',207-362 <SZO>
A:Cross-references: GB:M12678
A:Note: this allele is designated B*27052 (formerly 27W)
R:Vilches, C.
submitted to the EMBL Data Library, June 1993
A:Reference number: S34180
A:Accession: S34180
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-100,'S','102-137','D','139','Y','141-175','E',177-362 <VIL>
A:Cross-references: EMBL:X73578
A:Note: this allele is designated B*2706
R:D'Amato, M.; Sorrentino, R.
submitted to the EMBL Data Library, May 1994
A:Description: Identification of a novel HLA-B*27 subtype by restriction analysis of a c
A:Reference number: S44942
A:Accession: S44942
A:Molecule type: mRNA
A:Residues: 1-139,'H','141-362 <DAN>
A:Cross-references: EMBL:Z33453; NID:g486652; PIDN:CAA83876.1; PID:g486653
R:Ezquerria, A.; Bragado, R.; Vega, M.A.; Strominger, J.L.; Woody, J.; Lopez de Castro, J
Biochemistry 24, 1733-1741, 1985
A:Title: Primary structure of papain-solubilized human histocompatibility antigen HLA-B*2
A:Reference number: A30493; MUID:85226361
A:Accession: A30493
A:Molecule type: protein
A:Residues: 25-265,'E',267-295 <EZO>
R:Vega, M.A.; Ezquerria, A.; Rojo, S.; Aparicio, P.; Bragado, R.; Lopez de Castro, J.A.
Proc. Natl. Acad. Sci. U.S.A. 82, 7394-7398, 1985
A:Title: Structural analysis of an HLA-B*27 functional variant: identification of residue
A:Reference number: A94070; MUID:86042671
A:Accession: B24741
A:Molecule type: protein
A:Residues: 86-100,'N','102-103','IA',106-107;171-181 <VEG>
R:Coppin, H.L.; McDevitt, H.O.
J. Immunol. 137, 2168-2172, 1986
A:Title: Absence of polymorphism between HLA-B*27 genomic exon sequences isolated from no
A:Reference number: 135965; MUID:87009855
A:Accession: 135965
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 25-298 <RES>
A:Cross-references: GB:M14013; NID:g187743; PIDN:AAA59643.1; PID:g187744
R:Blasczyk, R.; Weber, M.; Salama, A.
submitted to the EMBL Data Library, January 1995
A:Reference number: S52291
A:Accession: S52291
A:Molecule type: DNA
A:Residues: 116-192 <BLA>
A:Cross-references: EMBL:X83737
A:Comment: This allele for HLA-B correlates with the development of ankylosing spondylit
C:Genetics:
A:Gene: GDB:HLA-B
A:Cross-references: GDB:120048; OMIM:142830
A:Map position: 6p21.3-6p21.3
A:Superfamily: class I histocompatibility antigen; immunoglobulin homology
A:Keywords: ankylosing spondylitis; duplication; glycoprotein; heterodimer; transmembran
F:1-24/Domain: signal sequence #status predicted <SIG>
F:25-362/Product: class I histocompatibility antigen HLA-B*27 alpha chain #status predict
F:25-307/Domain: extracellular #status predicted <EXT>
F:25-114/Domain: alpha-1 <EX1>
F:115-206/Domain: alpha-2 <EX2>

F:220-285/Domain: immunoglobulin homology <IMM>
F:308-331/Domain: transmembrane #status predicted <TMM>
F:332-362/Domain: intracellular #status predicted <INT>
F:110/Binding site: carbohydrate (Asn) (covalent) #status experimental
F:125-188,227-283/Disulfide bonds: #status experimental
Query Match 89.8%; Score 44; DB 1; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.22;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 REDLRILLRY 10
||||| |||||
Db 99 REDLRILLRY 108
RESULT 3
MHC class I histocompatibility antigen HLA-B*37 alpha chain precursor - human
C:Species: Homo sapiens (man)
C:Date: 16-Nov-1990 #sequence_revision 13-Jan-1993 #text_change 23-Jul-1999
C:Accession: C35997
R:Ennis, P.D.; Zemmour, J.; Salter, R.D.; Parham, P.
Proc. Natl. Acad. Sci. U.S.A. 87, 2833-2837, 1990
A:Title: Rapid cloning of HLA-A,B cDNA by using the polymerase chain reaction: freque
A:Reference number: A35997; MUID:90207291
A:Accession: C35997
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-362 <ENN>
A:Cross-references: GB:M32320; NID:g187792; PIDN:AAA36233.1; PID:g307224
C:Genetics:
A:Gene: GDB:HLA-B
A:Cross-references: GDB:120048; OMIM:142830
A:Map position: 6p21.3-6p21.3
A:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: transmembrane protein
F:220-285/Domain: immunoglobulin homology <IMM>
Query Match 89.8%; Score 44; DB 2; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.22;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 REDLRILLRY 10
||||| |||||
Db 99 REDLRILLRY 108
RESULT 4
MHC class I histocompatibility antigen HLA-B*47 precursor - human
C:Species: Homo sapiens (man)
C:Date: 07-Jun-1996 #sequence_revision 07-Jun-1996 #text_change 23-Jul-1999
C:Accession: I68724
R:Zemmour, J.; Ennis, P.D.; Parham, P.; Dupont, B.
Immunogenetics 27, 281-287, 1988
A:Title: Comparison of the structure of HLA-B*47 to HLA-B13 and its relationship to 2
A:Reference number: I54442; MUID:88152906
A:Accession: I68724
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-362 <ZEM>
A:Cross-references: GB:M19756; NID:g184171; PIDN:AAA52664.1; PID:g386776
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
Query Match 89.8%; Score 44; DB 2; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.22;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 REDLRILLRY 10
||||| |||||

Db 99 REDLRLTRY 108

RESULT 5

137485 human lymphocyte antigen HLA-B27 - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 23-Jul-1999
C:Accession: 137485
R:Del Porto, P.; D'Amato, M.; Fiorillo, M.T.; Tuosto, L.; Piccolella, E.; Sorrentino, R.
J. Immunol. 153, 3093-3100, 1994
A:Title: Identification of a novel HLA-B27 subtype by restriction analysis of a cytotoxic T lymphocyte epitope
A:Reference number: 137485; MUID:94375872
A:Accession: 137485
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-362 <RES>
A:Cross-references: EMBL:Z33453; NID:9486652; PIDN:CA83876.1; PID:9486653
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 89.8%; Score 44; DB 2; Length 362;

Best Local Similarity 90.0%; Pred. No. 0.22; 1; Indels 0; Gaps 0;
Matches 9; Conservative 0; Mismatches 0;

Qy 1 REDLRLTRY 10

||||| |||||
Db 99 REDLRLTRY 108

RESULT 6

154289 MHC HLA-B27d - human
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 23-Jul-1999
C:Accession: 154289
R:Choo, S.Y.; St. John, T.; Orr, H.T.; Hansen, J.A.
Hum. Immunol. 21, 209-219, 1988
A:Title: Molecular analysis of the variant alloantigen HLA-B27d (HLA-B*2703) identifies
A:Reference number: 154289; MUID:88227491
A:Accession: 154289
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-362 <RES>
A:Cross-references: GB:M54883; NID:g187663; PIDN:AAA59616.1; PID:g187664
C:Genetics:
A:Gene: GDB:HLA-B
A:Cross-references: GDB:120048; OMIM:142830
A:Map position: 6p21.3-6p21.3
A:Introns: 25/1; 115/1; 207/1; 299/1; 338/1; 349/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 89.8%; Score 44; DB 2; Length 362;

Best Local Similarity 90.0%; Pred. No. 0.22; 1; Indels 0; Gaps 0;
Matches 9; Conservative 0; Mismatches 0;

Qy 1 REDLRLTRY 10

||||| |||||
Db 99 REDLRLTRY 108

RESULT 7

180174 Class I histocompatibility antigen - chimpanzee (fragment)
C:Species: Pan troglodytes (chimpanzee)
C:Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 23-Jul-1999
C:Accession: I80174
R:McAdam, S.N.; Boyson, J.E.; Liu, X.; Garber, T.L.; Hughes, A.L.; Bontrop, R.E.; Watkins
Proc. Natl. Acad. Sci. U.S.A. 91, 5893-5897, 1994
A:Title: A uniquely high level of recombination at the HLA-B locus.
A:Reference number: 159308; MUID:94286544
A:Accession: I80174

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-137 <RES>

A:Cross-references: EMBL:U05585; NID:g454787; PIDN:AAA50188.1; PID:g454788
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 137;

Best Local Similarity 80.0%; Pred. No. 0.82; 1; Indels 0; Gaps 0;
Matches 8; Conservative 1; Mismatches 1;

Qy 1 REDLRLTRY 10

||||| |||||
Db 40 RENLRLTRY 49

RESULT 8

138509 MHC class I histocompatibility antigen - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 23-Jul-1999
C:Accession: 138509
R:Cereb, N.; Choi, J.W.; Riu, K.Z.; Yang, S.Y.
Tissue Antigens 44, 271-273, 1994
A:Title: HLA-B*5105, a newly identified B51 IEF variant.
A:Reference number: 138509; MUID:95176331
A:Accession: 138509
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-273 <RES>
A:Cross-references: EMBL:U06697; NID:g469544; PIDN:AAA92997.1; PID:g469545
C:Genetics:
A:Gene: GDB:HLA-B
A:Cross-references: GDB:120048; OMIM:142830
A:Map position: 6p21.3-6p21.3
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 273;

Best Local Similarity 80.0%; Pred. No. 1.7; 1; Indels 0; Gaps 0;
Matches 8; Conservative 1; Mismatches 1;

Qy 1 REDLRLTRY 10

||||| |||||
Db 74 RENLRLTRY 83

RESULT 9

154463 MHC HLA-B38 chain - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 07-Jun-1996 #sequence_revision 07-Jun-1996 #text_change 23-Jul-1999
C:Accession: 154463
R:Mueller, C.A.; Engler-Blum, G.; Gekeler, V.; Steiert, I.; Weiss, E.; Schmidt, H.
Immunogenetics 30, 200-207, 1989
A:Title: Genetic and serological heterogeneity of the supertypic HLA-B locus specific
A:Reference number: 154463; MUID:89379286
A:Accession: 154463
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-274 <RES>
A:Cross-references: GB:M29864; NID:g187674; PIDN:AAA36222.1; PID:g187675
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 274;

Best Local Similarity 80.0%; Pred. No. 1.7; 1; Indels 0; Gaps 0;
Matches 8; Conservative 1; Mismatches 1;

Qy 1 REDLRLTRY 10

||||| |||||
Db 75 RENLRLTRY 84

```
RESULT 10
class I histocompatibility antigen - pygmy chimpanzee (fragment)
C:Species: Pan paniscus (pygmy chimpanzee, bonobo)
C>Date: 31-May-1996 #sequence_revision 31-May-1996 #text_change 23-Jul-1999
C:Accession: I59308
R:McAdam, S.N.; Boyson, J.E.; Liu, X.; Garber, T.L.; Hughes, A.L.; Bontrop, R.E.; Watkin
Proc. Natl. Acad. Sci. U.S.A. 91, 5893-5897, 1994
A:Title: A uniquely high level of recombination at the HLA-B locus.
A:Reference number: I59308; MUID:94286544
A:Accession: I59308
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-354 <RES>
A:Cross-references: EMBL:U05575; NID:g454767; PIDN:AAA50178.1; PID:g454768
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 354;
Best Local Similarity 80.0%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
Db 91 RENLRALRY 100

RESULT 11
class I histocompatibility antigen - chimpanzee (fragment)
C:Species: Pan troglodytes (chimpanzee)
C>Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 23-Jul-1999
C:Accession: I80168
R:McAdam, S.N.; Boyson, J.E.; Liu, X.; Garber, T.L.; Hughes, A.L.; Bontrop, R.E.; Watkin
Proc. Natl. Acad. Sci. U.S.A. 91, 5893-5897, 1994
A:Title: A uniquely high level of recombination at the HLA-B locus.
A:Reference number: I59308; MUID:94286544
A:Accession: I80168
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-354 <RES>
A:Cross-references: EMBL:U05579; NID:g454775; PIDN:AAA50182.1; PID:g454776
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 354;
Best Local Similarity 80.0%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
Db 91 RENLRALRY 100

RESULT 12
class I histocompatibility antigen - pygmy chimpanzee (fragment)
C:Species: Pan paniscus (pygmy chimpanzee, bonobo)
C>Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 23-Jul-1999
C:Accession: I80167
R:McAdam, S.N.; Boyson, J.E.; Liu, X.; Garber, T.L.; Hughes, A.L.; Bontrop, R.E.; Watkin
Proc. Natl. Acad. Sci. U.S.A. 91, 5893-5897, 1994
A:Title: A uniquely high level of recombination at the HLA-B locus.
A:Reference number: I59308; MUID:94286544
A:Accession: I80167
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-354 <RES>
A:Cross-references: EMBL:U05578; NID:g454773; PIDN:AAA50181.1; PID:g454774
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
```

```
Query Match 79.6%; Score 39; DB 2; Length 354;
Best Local Similarity 80.0%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
Db 91 RENLRALRY 100

RESULT 13
class I histocompatibility antigen - chimpanzee (fragment)
C:Species: Pan troglodytes (chimpanzee)
C>Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 23-Jul-1999
C:Accession: I80169
R:McAdam, S.N.; Boyson, J.E.; Liu, X.; Garber, T.L.; Hughes, A.L.; Bontrop, R.E.; Wat
Proc. Natl. Acad. Sci. U.S.A. 91, 5893-5897, 1994
A:Title: A uniquely high level of recombination at the HLA-B locus.
A:Reference number: I59308; MUID:94286544
A:Accession: I80169
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-355 <RES>
A:Cross-references: EMBL:U05580; NID:g454777; PIDN:AAA50183.1; PID:g454778
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 355;
Best Local Similarity 80.0%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
Db 91 RENLRALRY 100

RESULT 14
class I histocompatibility antigen - chimpanzee (fragment)
C:Species: Pan troglodytes (chimpanzee)
C>Date: 24-May-1996 #sequence_revision 24-May-1996 #text_change 23-Jul-1999
C:Accession: I80171
R:McAdam, S.N.; Boyson, J.E.; Liu, X.; Garber, T.L.; Hughes, A.L.; Bontrop, R.E.; Wat
Proc. Natl. Acad. Sci. U.S.A. 91, 5893-5897, 1994
A:Title: A uniquely high level of recombination at the HLA-B locus.
A:Reference number: I59308; MUID:94286544
A:Accession: I80171
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: mRNA
A:Residues: 1-355 <RES>
A:Cross-references: EMBL:U05582; NID:g454781; PIDN:AAA50185.1; PID:g454782
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 79.6%; Score 39; DB 2; Length 355;
Best Local Similarity 80.0%; Pred. No. 2.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
Db 91 RENLRALRY 100

RESULT 15
MHC class I histocompatibility antigen HLA alpha chain precursor (clone pHLA 12.4) -
HLH12
C:Species: Homo sapiens (man)
C>Date: 05-Apr-1983 #sequence_revision 05-Apr-1983 #text_change 22-Jun-1999
C:Accession: A02189
R:Malissen, M.; Malissen, B.; Jordan, B.R.
Proc. Natl. Acad. Sci. U.S.A. 79, 893-897, 1982
A:Title: Exon/intron organization and complete nucleotide sequence of an HLA gene.
A:Reference number: A02189; MUID:82151002
```

A:Accession: A02189
A:Molecule type: DNA
A:Residues: 1-359 <MAL>
A:Cross-references: GB:J00191; GB:V00526; NID:g187600; PIDN:AAA36218.1; PID:g386873
C:Comment: The seven exons correspond approximately to the domain structure of this chain
C:Genetics:
A:Map position: 6p21.3
A:Introns: 22/1; 112/1; 204/1; 296/1; 335/1; 346/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: duplication; glycoprotein; heterodimer; transmembrane protein; transplantable
F:1-21/Domain: signal sequence #status predicted <SIG>
F:22-359/Product: class I histocompatibility antigen HLA alpha chain #status predicted
F:22-304/Domain: extracellular #status predicted <EXT>
F:22-111/Domain: alpha-1 <EX1>
F:112-203/Domain: alpha-2 <EX2>
F:217-282/Domain: immunoglobulin homology <IMM>
F:305-329/Domain: transmembrane #status predicted <TM>
F:335-359/Domain: intracellular #status predicted <INT>
F:107/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:224-280/Disulfide bonds: #status predicted

Query Match 79.6%; Score 39; DB 1; Length 359;
Best Local Similarity 80.0%; Pred.No. 2.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDLRILRY 10
 ||| |||
Db 96 RENLRILRY 105

Search completed: February 7, 2000, 11:54:25
Job time: 24335 sec

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DB 75 REDLRLTRY 84
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RESULT 2

ID IB14_HUMAN STANDARD: PRT: 361 AA.
AC P03989;
DT 23-OCT-1986 (Rel. 02, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-27 ALPHA CHAIN PRECURSOR.
GN HLA-B OR HLAB.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 86138405.
RA WEISS E.H., KUON W., DOERNER C., LANG M., RIETHMUELLER G.;
RT "Organization, sequence and expression of the HLA-B27 gene: a
RT molecular approach to analyze HLA and disease associations.";
RL Immunobiology 170:367-380(1985).
RN [2]
RP SEQUENCE OF 25-361 FROM N.A.
RX MEDLINE; 86149317.
RA SZOETS H., RIETHMUELLER G., WEISS E., MEO T.;
RT "Complete sequence of HLA-B27 CDNA identified through the
RT characterization of structural markers unique to the HLA-A, -B, and
RT -C allelic series.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:1428-1432(1986).
RN [3]
RP SEQUENCE OF 25-295.
RX MEDLINE; 85226361.
RA EZQUERRA A., BRAGADO R., VEGA M.A., STROMINGER J.L., WOODY J.,
RA LOPEZ DE CASTRO J.A.;
RT "Primary structure of papain-solubilized human histocompatibility
RT antigen HLA-B27.";
RL Biochemistry 24:1733-1741(1985).
RN [4]
RP X-RAY CRYSTALLOGRAPHY (2.1 ANGSTROMS) OF 25-300.
RX MEDLINE; 92405152.
RA MADDEN D.R., GORGA J.C., STROMINGER J.L., WILEY D.C.;
RT "The three-dimensional structure of HLA-B27 at 2.1-A resolution
RT suggests a general mechanism for tight peptide binding to MHC.";
RL Cell 70:1035-1048(1992).
RN [5]
RP X-RAY CRYSTALLOGRAPHY.
RX MEDLINE; 92018187.
RA MADDEN D.R., GORGA J.C., STROMINGER J.L., WILEY D.C.;
RT "The structure of HLA-B27 reveals nonamer self-peptides bound in an
RT extended conformation.";
RL Nature 353:321-325(1991).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC -!- DISEASE: THIS PROTEIN CORRELATES WITH THE DEVELOPMENT OF
CC ANKYLOSING SPONDYLITIS.
CC -----
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CC -----
CC EMBL; X03945; CAA27578.1; ALT_TERM.
CC PIR; A25128; HLHUB2.
CC PIR; S07441; S07441.
CC PDB; 1HSA; 15-OCT-92.
CC MIN; 142830; -.

DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; Ig; 1.
DR PFAM; PF00129; MHC_I; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; 3D-structure.
FT SIGNAL 1 24
FT CHAIN 25 361
FT FT
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT TRANSMEM 309 332
FT DOMAIN 333 361
FT CARBOHYD 110 110
FT DISULFID 125 188
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FT DISULFID 206 206
FT CONFLICT 266 266
FT STRAND 27 38
FT TURN 39 41
FT STRAND 42 52
FT TURN 53 54
FT STRAND 55 61
FT TURN 62 63
FT STRAND 70 71
FT HELIX 74 76
FT TURN 77 78
FT HELIX 81 108
FT TURN 109 110
FT TURN 113 114
FT STRAND 118 127
FT TURN 129 130
FT STRAND 133 142
FT TURN 143 144
FT STRAND 145 150
FT TURN 152 153
FT STRAND 157 159
FT HELIX 162 173
FT TURN 174 175
FT HELIX 176 185
FT TURN 186 186
FT HELIX 187 198
FT TURN 199 199
FT HELIX 200 203
FT TURN 204 204
FT STRAND 207 207
FT STRAND 210 217
FT STRAND 222 233
FT STRAND 238 243
FT TURN 244 245
FT STRAND 246 247
FT HELIX 249 251
FT STRAND 253 254
FT STRAND 258 259
FT STRAND 265 274
FT TURN 275 276
FT HELIX 278 280
FT STRAND 281 286
FT TURN 288 289
FT STRAND 294 296
SO SEQUENCE 361 AA; 40464 MW; 802130D5 CRC32;

Query Match 89.8%; Score 44; DB 1; Length 361;
Best Local Similarity 90.0%; Pred. No. 0.12;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRLTRY 10
| | | | |
DB 99 REDLRLTRY 108

RESULT 3

1B16_HUMAN
ID 1B16_HUMAN STANDARD; PRT; 362 AA.
AC P19373;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 01-FEB-1995 (Rel. 31, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-27 B*2703 ALPHA CHAIN
DE PRECURSOR (B-27D).
GN HLA-B OR HLAB.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
[1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 88227491.
RA CHOO S.Y., ST JOHN T., ORR H.T., HANSEN J.A.;
RT "Molecular analysis of the variant allc antigen HLA-B*2703 (HLA-B*2703)
RT identifies a unique single amino acid substitution.";
RL Hum. Immunol. 21:209-219(1988).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M54883; AAA59616.1; -.
DR HSP; P03989; 1HSA.
DR MIM; 142830; -.
DR PFAM; PS00290; IG_MHC; 1.
DR PROSITE; PF00047; 1g; 1.
DR PFAM; PF00129; MHC_I; 1.
DR MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 362
FT FT
FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT B-27 B*2703 ALPHA CHAIN.
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT TRANSMEM 309 332
FT DOMAIN 333 362
FT CARBOHYD 110 110
FT DISULFID 125 188
FT DISULFID 227 283
SQ SEQUENCE 362 AA; 40402 MW; 7261C3AB CRC32;
Query Match 89.8%; Score 44; DB 1; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.12;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 REDRLRLRY 10
Db 99 REDRLRLRY 108
RESULT 4
1B18_HUMAN STANDARD; PRT; 362 AA.
ID 1B18_HUMAN
AC P10318;
DT 01-MAR-1989 (Rel. 10, Created)
DT 01-MAR-1989 (Rel. 10, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-27 B*2705 ALPHA CHAIN
DE PRECURSOR (B-27W) (B27.1).
GN HLA-B OR HLAB.

OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
[1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 86220133.
RA SEEMANN G.H.A., REIN R.S., BROWN C.S., PLOEGH H.L.;
RT "Gene conversion-like mechanisms may generate polymorphism in human
RT class I genes.";
RL EMBO J. 5:547-552(1986).
[2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 86138405.
RA WEISS E.H., KUON W., DOERNER C., LANG M., RIETHMUELLER G.;
RT "Organizational, sequence and expression of the HLA-B*27 gene: a
RT molecular approach to analyze HLA and disease associations.";
RL Immunobiology 170:367-380(1985).
[3]
RP 3D-STRUCTURE MODELING OF 115-206.
RX MEDLINE; 95148615.
RA ROGNAN D., SCAPOZZA L., FOLKERS G., DASER A.;
RT "Rational design of nonnatural peptides as high-affinity ligands for
RT the HLA-B*2705 human leukocyte antigen.";
RL Proc. Natl. Acad. Sci. U.S.A. 92:753-757(1995).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
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CC -----
DR EMBL; X03665; CAA27302.1; -.
DR EMBL; X03666; CAA27302.1; JOINED.
DR EMBL; M12967; AAA36221.1; -.
DR PIR; A25092; HLHUBW.
DR PDB; 1ROG; 30-SEP-94.
DR PDB; 1ROH; 30-SEP-94.
DR PDB; 1ROI; 30-SEP-94.
DR PDB; 1ROJ; 30-SEP-94.
DR PDB; 1ROK; 30-SEP-94.
DR PDB; 1ROL; 30-SEP-94.
DR MIM; 142830; -.
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; 1g; 1.
DR PFAM; PF00129; MHC_I; 1.
DR MHC I; Transmembrane; Glycoprotein; Signal; 3D-structure.
FT SIGNAL 1 24
FT CHAIN 25 362
FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT B-27 B*2705 ALPHA CHAIN.
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT TRANSMEM 309 332
FT DOMAIN 333 362
FT CARBOHYD 110 110
FT DISULFID 125 188
FT DISULFID 227 283
SQ SEQUENCE 362 AA; 40428 MW; 73243566 CRC32;
Query Match 89.8%; Score 44; DB 1; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.12;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 REDRLRLRY 10
Db 99 REDRLRLRY 108
RESULT 4
1B18_HUMAN STANDARD; PRT; 362 AA.
ID 1B18_HUMAN
AC P10318;
DT 01-MAR-1989 (Rel. 10, Created)
DT 01-MAR-1989 (Rel. 10, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-27 B*2705 ALPHA CHAIN
DE PRECURSOR (B-27W) (B27.1).
GN HLA-B OR HLAB.

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Db 99 REDLRLLY 108

RESULT 5
ID 1B29_HUMAN STANDARD; PRT; 362 AA.
AC P18463;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-37 B*3701 ALPHA CHAIN
DE PRECURSOR.
GN HLA-B OR HLAB.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RA ZEMMOUR J., ENNIS P.D., PARHAM P., DUPONT B.;
RT "Comparison of the structure of HLA-B*47 to HLA-B13 and its
RT relationship to 21-hydroxylase deficiency.";
RL Immunogenetics 27:281-287(1988).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC
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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M19756; AAA52664.1; -
DR HSSP; P03989; ILSA.
DR MIM; 142830; -
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; Ig; 1.
DR PFAM; PF00129; MHC_I; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 362
FT FT
FT DOMAIN 25 114 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT DOMAIN 115 206 B-47 B*4701 ALPHA CHAIN.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-1.
FT DOMAIN 299 309 EXTRACELLULAR ALPHA-2.
FT TRANSMEM 310 333 EXTRACELLULAR ALPHA-3.
FT DOMAIN 333 362 CONNECTING PEPTIDE.
FT CARBOHYD 110 110 CYTOPLASMIC TAIL.
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
SQ SEQUENCE 362 AA; 40571 MW; 67734C1E CRC32;

Query Match 89.8%; Score 44; DB 1; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.12;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRLLY 10
Db 99 REDLRLLY 108

RESULT 7
ID 1B01_PANTR STANDARD; PRT; 359 AA.
AC P13750;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE CHLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-1 ALPHA CHAIN PRECURSOR
DE (FRAGMENT).
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Pan.
RN [1]
RP SEQUENCE FROM N.A.

```

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RX MEDLINE; 89030641.
RA MAYER W.E., JONKER M., KLEIN D., IVANYI P., VAN SEVENTER G.,
RT KLEIN J.;
RT "Nucleotide sequences of chimpanzee MHC class I alleles: evidence for
RL trans-species mode of evolution.";
RL EMBO J. 7:2765-2774(1988).
RN (2)
RP REVISIONS.
RA MAYER W.;
RL Submitted (FEB-1989) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
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CC -----
CC EMBL; X60255; CAA42807.1; -.
CC PIR; JH0539; JH0539.
CC HSP; P03989; IJSA.
CC PROSITE; PS00290; IG_MHC; 1.
CC PFAM; PF00047; Ig; 1.
CC MHC I; Transmembrane; Glycoprotein; Signal.
CC SIGNAL 1 24
CC CHAIN 25 362
CC BY SIMILARITY.
CC CLASS I HISTOCOMPATIBILITY ANTIGEN,
CC GOGO-B0101 ALPHA CHAIN.
CC EXTRACELLULAR ALPHA-1.
CC EXTRACELLULAR ALPHA-2.
CC EXTRACELLULAR ALPHA-3.
CC CONNECTING PEPTIDE.
CC TRANSMEM 309 332
CC DOMAIN 333 362
CC DISULFID 125 188
CC DISULFID 227 283
CC CARBOHYD 110 110
CC BY SIMILARITY.
CC SEQUENCE 362 AA; 40170 MW; 2E33E2B8 CRC32;

Query Match 79.6%; Score 39; DB 1; Length 362;
Best Local Similarity 80.0%; Pred. No. 1.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILRY 10
Db 95 RENDLRALRY 108

RESULT 9
1B01_GORGO
ID 1B01_GORGO STANDARD; PRT; 362 AA.
AC P30379;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DE CLASS I HISTOCOMPATIBILITY ANTIGEN, GOGO-B0102 ALPHA CHAIN PRECURSOR.
OS Gorilla gorilla gorilla (Lowland gorilla).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Gorilla.
RN (1)
RP SEQUENCE FROM N.A.
RX MEDLINE; 92078860.
RA LAWOR D.A., WARREN E., TAYLOR P., PARHAM P.;
RT "Gorilla class I major histocompatibility complex alleles: comparison
RL to human and chimpanzee class I.";
RL J. Exp. Med. 174:1491-1509(1991).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
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CC -----
CC EMBL; X13115; CAA31507.1; -.
CC PIR; S03537; S03537.
CC HSP; P03989; IJSA.
CC PROSITE; PS00290; IG_MHC; 1.
CC PFAM; PF00047; Ig; 1.
CC MHC I; Transmembrane; Glycoprotein; Signal.
CC SIGNAL 1 20
CC CHAIN 21 359
CC BY SIMILARITY.
CC CLASS I HISTOCOMPATIBILITY ANTIGEN,
CC B-1 ALPHA CHAIN.
CC EXTRACELLULAR ALPHA-1.
CC EXTRACELLULAR ALPHA-2.
CC EXTRACELLULAR ALPHA-3.
CC CONNECTING PEPTIDE.
CC TRANSMEM 306 329
CC DOMAIN 330 359
CC DISULFID 121 184
CC DISULFID 223 279
CC CARBOHYD 106 106
CC BY SIMILARITY.
CC SEQUENCE 359 AA; 40173 MW; 5395FFC9 CRC32;

Query Match 79.6%; Score 39; DB 1; Length 359;
Best Local Similarity 80.0%; Pred. No. 1.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILRY 10
Db 95 RENDLRALRY 104

RESULT 8
1B01_GORGO
ID 1B01_GORGO STANDARD; PRT; 362 AA.
AC P30379;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DE CLASS I HISTOCOMPATIBILITY ANTIGEN, GOGO-B0101 ALPHA CHAIN PRECURSOR.
OS Gorilla gorilla gorilla (Lowland gorilla).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Gorilla.
RN (1)
RP SEQUENCE FROM N.A.
RX MEDLINE; 92078860.
RA LAWOR D.A., WARREN E., TAYLOR P., PARHAM P.;
RT "Gorilla class I major histocompatibility complex alleles: comparison
RL to human and chimpanzee class I.";
RL J. Exp. Med. 174:1491-1509(1991).

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CC -----

CC EMBL; M22319; AAA36232.1; -;
DR EMBL; M22792; AAA59620.1; ALT_SEQ.
DR EMBL; M22786; AAA59620.1; JOINED.
DR EMBL; M22787; AAA59620.1; JOINED.
DR EMBL; M22788; AAA59620.1; JOINED.
DR EMBL; M22789; AAA59620.1; JOINED.
DR EMBL; M22790; AAA59620.1; JOINED.
DR EMBL; M22791; AAA59620.1; JOINED.
DR EMBL; M22792; AAA59620.1; JOINED.
DR EMBL; L41087; AAA64513.1; -;
DR EMBL; L41086; AAA64513.1; JOINED.
DR PIR; A30345; A30345.
DR PIR; A30548; A30548.
DR HSP; P30491; IAIM.
DR MIM; 142830; -;
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; IG; 1.
DR PFAM; PF00129; MHC_I; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 362
FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT B-51(B-5) B*5101 ALPHA CHAIN.
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT DOMAIN 309 332
FT TRANSMEM 333 362
FT DOMAIN 333 362
FT CARBOHYD 110 110
FT BY SIMILARITY.
FT DISULFID 125 188
FT BY SIMILARITY.
FT DISULFID 227 283
SQ SEQUENCE 362 AA; 40566 MW; 4D846F30 CRC32;

Query Match 79.6%; Score 39; DB 1; Length 362;
Best Local Similarity 80.0%; Pred. No. 1.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLILLRY 10
II:III III
DB 99 RENLRALRY 108

RESULT 14
ID 1B52_HUMAN STANDARD; PRT; 362 AA.
AC P30489;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, B-51(B-5) B*5104 ALPHA CHAIN
DE PRECURSOR.
GN HLA-B OR HLAB.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 92269955.
RA BELICH M.P., MADRICAL J.A., HILDEBRAND W.H., ZEMMOUR J.,
RA WILLIAMS R.C., LUZ R., PETZL-ERLER M.L., PARHAM P.;
RT "Unusual HLA-B alleles in two tribes of Brazilian Indians."
RL Nature 357:326-329(1992).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
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CC EMBL; M22799; AAA59645.1; ALT_SEQ.
DR EMBL; M22793; AAA59645.1; JOINED.
DR EMBL; M22794; AAA59645.1; JOINED.

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CC EMBL; Z15143; CAA78849.1; -;
DR HSP; P30491; IAIM.
DR MIM; 142830; -;
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; IG; 1.
DR PFAM; PF00129; MHC_I; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 362
FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT B-51(B-5) B*5104 ALPHA CHAIN.
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT DOMAIN 309 332
FT TRANSMEM 333 362
FT DOMAIN 333 362
FT CARBOHYD 110 110
FT BY SIMILARITY.
FT DISULFID 125 188
FT BY SIMILARITY.
FT DISULFID 227 283
SQ SEQUENCE 362 AA; 40560 MW; F22F08AB CRC32;

Query Match 79.6%; Score 39; DB 1; Length 362;
Best Local Similarity 80.0%; Pred. No. 1.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLILLRY 10
II:III III
DB 99 RENLRALRY 108

RESULT 15
ID 1B53_HUMAN STANDARD; PRT; 362 AA.
AC P30490;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, BW-52(B-5) B*5201 ALPHA CHAIN
DE PRECURSOR.
GN HLA-B OR HLAB.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 89080265.
RA HAYASHI H., ENNIS P.D., ARIGA H., SALTER R.D., PARHAM P., KANO K.,
RA TAKIGUCHI M.;
RT "HLA-B51 and HLA-Bw52 differ by only two amino acids which are in the
RT helical region of the alpha 1 domain."
RL J. Immunol. 142:306-311(1989).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
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CC EMBL; M22799; AAA59645.1; ALT_SEQ.
DR EMBL; M22793; AAA59645.1; JOINED.
DR EMBL; M22794; AAA59645.1; JOINED.

DR EMBL; M22795; AAA59645.1; JOINED.
 DR EMBL; M22796; AAA59645.1; JOINED.
 DR EMBL; M22797; AAA59645.1; JOINED.
 DR EMBL; M22798; AAA59645.1; JOINED.
 DR PIR; B30345; B30345.
 DR PIR; B30548; B30548.
 DR HSP; P30491; IAIM.
 DR MIN; 142830; .
 DR PROSITE; PS00290; IG_MHC; 1.
 DR PFAM; PF00047; ig; 1.
 DR PFAM; PF00129; MHC_I; 1.
 KW MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 24
 FT CHAIN 25 362 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
 BW-52(B-5) B*5201 ALPHA CHAIN.
 FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 299 308 CONNECTING PEPTIDE.
 FT TRANSMEM 309 332 CYTOPLASMIC TAIL.
 FT DOMAIN 333 362 BY SIMILARITY.
 FT CARBOHYD 110 110 BY SIMILARITY.
 FT DISULFID 125 188 BY SIMILARITY.
 FT DISULFID 227 283 BY SIMILARITY.
 SQ SEQUENCE 362 AA; 40521 MW; 3B436FE8 CRC32;

Query Match 79.6%; Score 39; DB 1; Length 362;
 Best Local Similarity 80.0%; Pred. No. 1.2;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDLRILRY 10
 Db 99 RNLRLRY 108

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 Job time: 3782 sec

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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:41 ; Search time 209.03 seconds
(without alignments)
3.317 Million cell updates/sec

Title: US-08-653-294-14
Perfect score: 49
Sequence: 1 REDRLILLRY 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 225878 seqs, 69334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : SPTREMBL12:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	% Match	Query Length	DB ID	Description
1	44	89.8	39	7	019688 homo sapien
2	44	89.8	90	7	019193 homo sapien
3	44	89.8	181	7	062898 homo sapien
4	44	89.8	181	7	078138 homo sapien
5	44	89.8	181	7	078142 homo sapien
6	44	89.8	274	7	019692 homo sapien
7	44	89.8	322	7	019627 homo sapien
8	44	89.8	359	7	029934 homo sapien
9	44	89.8	362	7	029705 homo sapien
10	44	89.8	362	7	029846 homo sapien
11	44	89.8	362	7	078189 homo sapien
12	39	79.6	89	7	019569 homo sapien
13	39	79.6	90	7	046697 gorilla gor
14	39	79.6	133	7	019189 homo sapien
15	39	79.6	137	7	095533 pan troglod
16	39	79.6	138	7	078209 homo sapien
17	39	79.6	172	7	019770 homo sapien
18	39	79.6	172	7	019774 homo sapien
19	39	79.6	172	7	019775 homo sapien
20	39	79.6	172	7	019780 homo sapien

21	39	79.6	172	7	Q95364	homo sapien
22	39	79.6	172	7	019771	homo sapien
23	39	79.6	172	7	019772	homo sapien
24	39	79.6	172	7	019773	homo sapien
25	39	79.6	175	7	Q29694	homo sapien
26	39	79.6	180	7	019607	homo sapien
27	39	79.6	180	7	019608	homo sapien
28	39	79.6	180	7	019609	homo sapien
29	39	79.6	180	7	019610	homo sapien
30	39	79.6	180	7	019611	homo sapien
31	39	79.6	180	7	019612	homo sapien
32	39	79.6	180	7	019613	homo sapien
33	39	79.6	181	7	046703	homo sapien
34	39	79.6	181	7	062917	homo sapien
35	39	79.6	181	7	062892	homo sapien
36	39	79.6	181	7	062899	homo sapien
37	39	79.6	181	7	062920	homo sapien
38	39	79.6	181	7	062922	homo sapien
39	39	79.6	181	7	062923	homo sapien
40	39	79.6	181	7	019623	homo sapien
41	39	79.6	181	7	019747	homo sapien
42	39	79.6	181	7	Q29667	homo sapien
43	39	79.6	181	7	Q30198	homo sapien
44	39	79.6	181	7	Q29708	homo sapien
45	39	79.6	181	7	Q29724	homo sapien

ALIGNMENTS

RESULT 1
019688 PRELIMINARY; PRT; 39 AA.
AC 019688;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
DE HLA-B27 VARIANT EXON 2 (ALPHAL DOMAIN) (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RA BLASZYK R., WEBER M., SALAMA A.;
RL Submitted (JAN-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; X83727; CAA58698.1; -
DR PFAM; PF00129; MHC_I; 1.
KW MHC.
FT NON_TER 1 1
FT NON_TER 39 39
SQ SEQUENCE 39 AA; 4748 MW; 6F714D4C CRC32;

Query Match 89.8%; Score 44; DB 7; Length 39;
Best Local Similarity 90.0%; Pred. No. 0.07;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLILLRY 10
DB 24 REDRLILLRY 33

RESULT 2
019193 PRELIMINARY; PRT; 90 AA.
AC 019193;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
DE HISTOCOMPATIBILITY ANTIGEN ALPHA 1 DOMAIN (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE: 92337445.
 RA HIGGINS C.M., LUND T., SHIPLEY M.E., EBRINGER A.,
 RA SADOWSKA-WROBLEWSKA M., CRAIG R.K.;
 RT "Ankylosing spondylitis and HLA-B*27: restriction fragment length
 RT polymorphism and sequencing of an HLA-B*27 allele from a patient with
 RT ankylosing spondylitis."
 RL Ann. Rheum. Dis. 51:855-862(1992).
 DR EMBL: S39758; CAB27364.1; -.
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 FT NON_TER 1 1
 FT NON_TER 90 90
 SQ SEQUENCE 90 AA; 10571 MW; F22CCB4E CRC32;

Query Match 89.8%; Score 44; DB 7; Length 90;
 Best Local Similarity 90.0%; Pred. No. 0.17;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
 ||||| ||||
 Db 75 REDLRLLRY 84

RESULT 3
 O62898 PRELIMINARY; PRT; 181 AA.
 ID O62898
 AC O62898
 DT 01-AUG-1998 (TREMBLrel. 07, Created)
 DT 01-AUG-1998 (TREMBLrel. 07, Last sequence update)
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
 DE MHC CLASS I ANTIGEN (FRAGMENT).
 GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA KOSMAN C.A., HURLEY C.K.;
 RL Submitted (MAR-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF054012; AAC32563.1; -.
 DR EMBL: AF054011; AAC32563.1; JOINED.
 DR HSSP: P10318; IROG.
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 FT NON_TER 1 1
 FT NON_TER 181 181
 SQ SEQUENCE 181 AA; 21107 MW; D8E533DD CRC32;

Query Match 89.8%; Score 44; DB 7; Length 181;
 Best Local Similarity 90.0%; Pred. No. 0.34;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
 ||||| ||||
 Db 74 REDLRLLRY 83

RESULT 4
 O78138 PRELIMINARY; PRT; 181 AA.
 ID O78138
 AC O78138
 DT 01-NOV-1998 (TREMBLrel. 08, Created)
 DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
 DE MHC CLASS I ANTIGEN (FRAGMENT).
 GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.

OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA KOSMAN C.A., HURLEY C.K.;
 RT "Novel HLA Class I B Locus Alleles."
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF071770; AAC35939.1; -.
 DR EMBL: AF071769; AAC35939.1; JOINED.
 DR HSSP: P10318; IROG.
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 FT NON_TER 1 1
 FT NON_TER 181 181
 SQ SEQUENCE 181 AA; 21103 MW; 8CF468CF CRC32;

Query Match 89.8%; Score 44; DB 7; Length 181;
 Best Local Similarity 90.0%; Pred. No. 0.34;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
 ||||| ||||
 Db 74 REDLRLLRY 83

RESULT 5
 O78142 PRELIMINARY; PRT; 181 AA.
 ID O78142
 AC O78142
 DT 01-NOV-1998 (TREMBLrel. 08, Created)
 DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
 DE MHC CLASS I ANTIGEN (FRAGMENT).
 GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA STEINER N.K., HURLEY C.K., KOESTER R.P.;
 RT "Novel HLA-B allele."
 RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF072764; AAC25779.1; -.
 DR EMBL: AF072763; AAC25779.1; JOINED.
 DR HSSP: P10318; IROG.
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 FT NON_TER 1 1
 FT NON_TER 181 181
 SQ SEQUENCE 181 AA; 21079 MW; 24949B0F CRC32;

Query Match 89.8%; Score 44; DB 7; Length 181;
 Best Local Similarity 90.0%; Pred. No. 0.34;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDLRILLRY 10
 ||||| ||||
 Db 74 REDLRLLRY 83

RESULT 6
 O19692 PRELIMINARY; PRT; 274 AA.
 ID O19692
 AC O19692
 DT 01-JAN-1998 (TREMBLrel. 05, Created)
 DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
 DE MHC CLASS I HLA-B*27 M (FRAGMENT).
 GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.

```

RN RP SEQUENCE FROM N.A.
RX MEDLINE: 87009855.
RA COPPIN H.L., MCDEVITT H.O.;
RT "Absence of polymorphism between HLA-B27 genomic exon sequences
RT isolated from normal donors and ankylosing spondylitis patients.";
RL J. Immunol. 137:2168-2172(1986).
DR EMBL; M14013; AAA59643.1; -.
DR HSSP; P10318; IROG.
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; Ig; 1.
DR PFAM; PF00129; MHC_I; 1.
DR MHC.
KW NON_TER 1 1
FT NON_TER 274 274
SQ SEQUENCE 274 AA; 31659 MW; 9A74A6BA CRC32;

Query Match 89.8%; Score 44; DB 7; Length 274;
Best Local Similarity 90.0%; Pred. No. 0.53;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
Db 75 REDRLRLRY 84

RESULT 7
O19627 PRELIMINARY; PRT; 322 AA.
AC O19627;
DT 01-JAN-1998 (TREMBLrel. 05, Created)
DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
DE HLA-B37 (FRAGMENT).
GN B-3701.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN RP SEQUENCE FROM N.A.
RA HURLEY C.K., BEI M., RODRIGUEZ S., JOHNSON A.;
RL Submitted (JUN-1994) to the EMBL/GenBank/DBJ databases.
DR EMBL; U11267; AAA19927.1; -.
DR HSSP; P30685; IA9E.
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; Ig; 1.
DR PFAM; PF00129; MHC_I; 1.
DR MHC.
KW NON_TER 322 322
FT NON_TER 322 322
SQ SEQUENCE 322 AA; 36626 MW; DF3B7744 CRC32;

Query Match 89.8%; Score 44; DB 7; Length 322;
Best Local Similarity 90.0%; Pred. No. 0.62;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
Db 99 REDRLRLRY 108

RESULT 8
O29934 PRELIMINARY; PRT; 359 AA.
AC O29934;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
DE HLA-B27 (FRAGMENT).
GN HLA-B.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

[1]
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN RP SEQUENCE FROM N.A.
RX MEDLINE: 86149317.
RA SZOTS H., RIETHMULLER G., WEISS E., MEO T.;
RT "Complete sequence of HLA-B27 cDNA identified through the
RT characterization of structural markers unique to the HLA-A, -B, and -C
RT allelic series.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:1428-1432(1986).
DR EMBL; M12678; AAA59614.1; -.
DR HSSP; P10318; IROG.
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; Ig; 1.
DR PFAM; PF00129; MHC_I; 1.
DR MHC.
KW NON_TER 1 1
FT NON_TER 359 359
SQ SEQUENCE 359 AA; 40042 MW; 069F7E64 CRC32;

Query Match 89.8%; Score 44; DB 7; Length 359;
Best Local Similarity 90.0%; Pred. No. 0.69;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
Db 96 REDRLRLRY 105

RESULT 9
O29705 PRELIMINARY; PRT; 362 AA.
AC O29705;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
DE MHC CLASS I ANTIGEN HLA-B PRECURSOR.
GN HLA-B.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN RP SEQUENCE FROM N.A.
RA BALAS A., SANTOS S., VICARIO J.L.;
RL Submitted (JUL-1995) to the EMBL/GenBank/DBJ databases.
DR EMBL; U31971; AAA98506.1; -.
DR HSSP; P10318; IROG.
DR PROSITE; PS00290; IG_MHC; 1.
DR PFAM; PF00047; Ig; 1.
DR PFAM; PF00129; MHC_I; 1.
DR MHC.
KW SIGNAL 1 24
FT CHAIN 25 362
FT CHAIN 25 362
SQ SEQUENCE 362 AA; 40479 MW; 09C9D20A CRC32;

Query Match 89.8%; Score 44; DB 7; Length 362;
Best Local Similarity 90.0%; Pred. No. 0.7;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 REDRLRLRY 10
Db 99 REDRLRLRY 108

RESULT 10
O29846 PRELIMINARY; PRT; 362 AA.
AC O29846;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
DE HUMAN LYMPHOCYTE ANTIGEN HLA-B27.

```

OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=BLOOD;
 RX MEDLINE: 94375872.
 RA DEL PORTO P., D'AMATO M., FIORILLO M.T., TUOSTO L., PICCOLELLA E.,
 RA SORRENTINO R.;
 RT Identification of a novel HLA-B*27 subtype by restriction analysis of
 RT a cytotoxic gamma delta T cell clone.";
 RL J. Immunol. 153:3093-3100(1994).
 DR EMBL: Z33453; CAA83876.1; -
 DR HSSP: P10318; IROG.
 DR PROSITE: PS00290; IG_MHC; 1.
 DR PFAM: PF00047; Ig; 1.
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 SQ SEQUENCE 362 AA; 40450 MW; CCA23A50 CRC32;

Query Match 89.8%; Score 44; DB 7; Length 362;
 Best Local Similarity 90.0%; Pred. No. 0.7;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLLY 10
 ||||| ||||
 Db 99 REDRLRLLY 108

RESULT 11
 O78189 PRELIMINARY; PRT; 362 AA.
 AC O78189;
 DT 01-NOV-1998 (TEMBLrel. 08, Created)
 DT 01-NOV-1998 (TEMBLrel. 08, Last sequence update)
 DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
 DE MHC CLASS I ANTIGEN.
 GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA SEURYNCK K.L., BAXTER-LOWE L.A.;
 RT "B27052 W496D.";
 RL Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF026218; AAC42275.1; -
 DR HSSP: P10318; IROG.
 DR PROSITE: PS00290; IG_MHC; 1.
 DR PFAM: PF00047; Ig; 1.
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 SQ SEQUENCE 362 AA; 40486 MW; 2B0EF602 CRC32;

Query Match 89.8%; Score 44; DB 7; Length 362;
 Best Local Similarity 90.0%; Pred. No. 0.7;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLLY 10
 ||||| ||||
 Db 99 REDRLRLLY 108

RESULT 12
 O19569 PRELIMINARY; PRT; 89 AA.
 AC O19569;
 DT 01-JAN-1998 (TEMBLrel. 05, Created)
 DT 01-MAY-1999 (TEMBLrel. 10, Last sequence update)
 DT 01-MAY-1999 (TEMBLrel. 10, Last annotation update)
 DE MHC CLASS I ANTIGEN (FRAGMENT).

GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA CAO K., BURDETT L., ZHANG G., FERNANDEZ-VINA M.;
 RL Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF017320; AAB70286.2; -
 KW MHC.
 FT NON_TER 1
 FT NON_TER 89
 SQ SEQUENCE 89 AA; 10606 MW; 99D11089 CRC32;

Query Match 79.6%; Score 39; DB 7; Length 89;
 Best Local Similarity 80.0%; Pred. No. 1.6;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLLY 10
 ||||| ||||
 Db 74 REDRLRLLY 83

RESULT 13
 O46697 PRELIMINARY; PRT; 90 AA.
 AC O46697;
 DT 01-JUN-1998 (TEMBLrel. 06, Created)
 DT 01-JUN-1998 (TEMBLrel. 06, Last sequence update)
 DT 01-MAY-1999 (TEMBLrel. 10, Last annotation update)
 DE MHC CLASS I ANTIGEN HLA-H ORTHOLOG (FRAGMENT).
 GN HLA-H.
 OS Gorilla gorilla gorilla (Lowland gorilla).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Gorilla.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=SHAMBA;
 RA GRIMSLEY C., MATHER K.A., OBER C.;
 RL Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF022172; AAC99794.1; -
 DR PFAM: PF00129; MHC_I; 1.
 KW MHC.
 FT NON_TER 1
 FT NON_TER 90
 SQ SEQUENCE 90 AA; 10689 MW; 5E5F2495 CRC32;

Query Match 79.6%; Score 39; DB 7; Length 90;
 Best Local Similarity 80.0%; Pred. No. 1.6;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 REDRLRLLY 10
 ||||| ||||
 Db 75 REDRLRLLY 84

RESULT 14
 O19189 PRELIMINARY; PRT; 133 AA.
 AC O19189;
 DT 01-JAN-1998 (TEMBLrel. 05, Created)
 DT 01-JAN-1998 (TEMBLrel. 05, Last sequence update)
 DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
 DE MHC CLASS I HISTOCOMPATIBILITY ANTIGEN-B (HLA-B-27KSH) (FRAGMENT).
 GN HLA-B.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=LEUKOCYTE;

RA PETERSDORF E.;
 RL Submitted (DEC-1994) to the EMBL/GenBank/DBJ databases.
 DR EMBL; U18659; IAB60357.1; -;
 DR MIM; 142830; -;
 DR PFAM; PF00129; MHC_I; 1.
 KW MHC I.

FT NON_TER 1 1
 FT NON_TER 133 133
 SQ SEQUENCE 133 AA; 15491 MW; 3A3BC802 CRC32;

Query Match 79.6%; Score 39; DB 7; Length 133;
 Best Local Similarity 80.0%; Pred. No. 2.4;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 REDLRILLRY 10
 ||:|||||
 Db 27 RENLRALRY 36

RESULT 15

Q95533 PRELIMINARY; PRT; 137 AA.
 AC Q95533;
 DT 01-FEB-1997 (TREMBlrel. 02, Created)
 DT 01-FEB-1997 (TREMBlrel. 02, Last sequence update)
 DT 01-NOV-1998 (TREMBlrel. 08, Last annotation update)
 DE CLASS I HISTOCOMPATIBILITY ANTIGEN (FRAGMENT).
 GN HLA-B.
 OS Pan troglodytes (Chimpanzee).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Primates; Catarrhini; Hominidae; Pan.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-WODKA;
 RX MEDLINE; 94286544.
 RA MCADAM S.N., BOYSON J.E., LIU X., GARBER T.L., HUGHES A.L.,
 RA BONTROP R.E., WATKINS D.I.;
 RT "A uniquely high level of recombination at the HLA-B locus."
 RL Proc. Natl. Acad. Sci. U.S.A. 91:5893-5897(1994).
 DR EMBL; U05585; AAA50188.1; -;
 DR PFAM; PF00129; MHC_I; 1.
 KW MHC.
 FT NON_TER 1 1
 FT NON_TER 137 137
 SQ SEQUENCE 137 AA; 15922 MW; B316D3BC CRC32;

Query Match 79.6%; Score 39; DB 7; Length 137;
 Best Local Similarity 80.0%; Pred. No. 2.4;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 1 REDLRILLRY 10
 ||:|||||
 Db 40 RENLRLLRY 49

Search completed: February 8, 2000, 13:17:41
 Job time: 32490 sec

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OM of: US-08-653-294-14 to: GenEmbl.* out_format : pfs

Date: Feb 8, 2000 4:40 PM

About: Results were produced by the GenCore software, version 4.5.
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Command line parameters:

-MODEL=frame+ p2n.model -DEV=xlp
-O=/cgnl_1/USPTO.spool/US08653294/runat_04022000_160701_15779/app_query.fasta.1
-DB=GenEmbl -QFMT=fastap -SUFFIX=rge -GAPOP=12.000 -GAPEXT=4.000
-MINMATCH=0.100 -LOOPEXT=0.000 -LOOPEXT=0.000 -GAPOP=4.500
-GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -GAPOP=6.000
-GAPEXT=7.000 -XGAPOP=10.000 -XGAPEXT=0.500 -DELOP=6.000
-DELEXT=7.000 -START=1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOALIGN=200 -THR SCORE=pct -ALIGN=15 -MODE=LOCAL
-OUTFMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=100000 -USER=US08653294
-NCPU=6 -ICPU=3 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-08-653-294-14

Query length: 10

Database: GenEmbl.*

Database sequences: 821193

Database length: -1518192014

Search time (sec): 11370.480000

score_list:

Sequence	Strd Orig	ZScore	Escore	Len	Documentation
gb_pr1:HSAB27V2	+	44.00	170.85	195	! X83727 H.sapiens HLA-B27 variant
gb_pr3:HSAB27HLA1	+	44.00	168.07	270	! AF072763 Homo sapiens MHC class
gb_pr3:HSB1524V1	+	44.00	168.07	270	! AF054011 Homo sapiens isolate C
gb_pr3:HSBCHLBC1	+	44.00	168.07	270	! AF071769 Homo sapiens MHC class
gb_pr4:HS27052B1	+	44.00	168.07	270	! AF102563 Homo sapiens MHC class
gb_pr4:HS14AWBY2	+	44.00	168.07	270	! AF110257 Homo sapiens MHC class
gb_pr2:HUMB27052G	+	44.00	162.07	546	! L76095 Homo sapiens MHC class I
gb_pr1:HUMHBM2	+	44.00	158.58	822	! M14013 Human MHC class I HLA-B2
gb_pr1:HSU11267	+	44.00	157.19	968	! U11267 Human HLA-B37 (B-3701) m
gb_pr1:HUMHMC	+	44.00	156.77	1017	! M62852 Human MHC class I HLA-E
gb_pr1:HSAB27V2	+	44.00	156.69	1026	! A28264 H.sapiens mRNA for HLA-
gb_pr1:HSAB27HLA1	+	44.00	156.69	1026	! E01342 cDNA encoding C-termina
gb_pr1:HUMHBM2	+	44.00	156.22	1112	! L20086 Human MHC class I (HLA-
gb_pr2:HSU31971	+	44.00	156.19	1089	! U31971 Human MHC class I antiq
gb_pr3:AF026218	+	44.00	156.15	1089	! AF026218 Homo sapiens MHC clas
gb_pr1:HUMHBM2	+	44.00	156.15	1093	! M19756 Human MHC class I HLA-E
gb_pr1:HUMHBM2	+	44.00	156.15	1093	! M32320 Human MHC HLA protein,
gb_pr1:HSAB27V2	+	44.00	155.07	1241	! X03665 Human class I MHC gene
gb_pr1:HSAB27V2	+	44.00	154.86	1272	! Z33453 H.sapiens mRNA for huma
gb_pr1:HUMHBM2	+	44.00	153.42	1507	! M12678 Human HLA-B27 mRNA, con
gb_pr1:HUMHBM2	+	44.00	145.88	3649	! M54883 Human MHC class I HLA-E
gb_pr1:HUMHBM2	+	44.00	145.06	4015	! E01341 Genomic DNA encoding H
gb_pr1:HUMHBM2	+	44.00	145.06	4016	! M12967 Human MHC class I HLA-E
gb_pr4:S39758	+	44.00	144.59	4242	! S39758 HLA-B27 (HLA-B*2705)-hl
gb_pr1:HSAB27V2	+	44.00	144.50	501	! X03945 Human gene for HLA-B27
gb_pr1:HSAB27V2	+	44.00	140.89	6553	! AR008238 Sequence 1 from paten
gb_pr1:HSAB27V2	+	42.00	106.88	624.87	! D90899 Synchocystis sp. PCC
gb_pr1:HSAB27V2	+	42.00	106.88	624.87	! M58352 Proteus mirabilis 60k-r
gb_pr1:HSAB27V2	+	40.00	114.59	232.40	! AC016616 Homo sapiens chromos
gb_pr1:HSAB27V2	+	40.00	108.29	871.32	! AC013786 Homo sapiens chromos
gb_pr1:HSAB27V2	+	40.00	101.85	1.2e+03	! AC006837 Arabidopsis thaliana
gb_pr1:HSAB27V2	+	40.00	101.66	1.2e+03	! AL021811 Arabidopsis thaliana
gb_pr1:HSAB27V2	+	40.00	100.29	1.5e+03	! AL050402 Human DNA sequence
gb_pr1:HSAB27V2	+	40.00	95.27	2.4e+03	! AC009409 Homo sapiens clone
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gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y08692 H.sapiens HLA-B gene, ex
gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y08693 H.sapiens HLA-B gene, ex
gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y08694 H.sapiens HLA-B gene, ex
gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y04020 Human cell line THAI DCH
gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y06111 Human cell line THAI DCH
gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y06113 Human cell line THAI DCH
gb_pr1:HSAB27V2	+	39.00	148.01	3.20	! Y06115 Human cell line THAI DCH

gb_pr2:HSAB27V2 + 39.00 148.01 3.20 250 ! U90422 Human cell line THAI
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gb_pr2:HSAB27V2 + 39.00 148.01 3.20 250 ! U90418 Human cell line THAI

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seq_documentation_block:

LOCUS HSLAB27V2 195 bp DNA PRI 31-JAN-1995
DEFINITION H.sapiens HLA-B27 variant gene (exon 2).

ACCESSION X83727

VERSION X83727.1 GI:663002

KEYWORDS HLA-B gene; human leukocyte antigen; major histocompatibility complex class I.

SOURCE human.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 195)

AUTHORS Blasczyk,R., Weber,M. and Salama,A.

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 195)

AUTHORS Blasczyk,R.

TITLE Direct Submission

JOURNAL Submitted (06-JAN-1995) R. Blasczyk, Bloodbank, Dept.of Intern.

Medicine, Div of Hematol. and Oncolog., Spandauer Damm 130, Univ.

Hosp. Rudolf Virchow, Freie Univ., D- 14050 Berlin, FRG

FEATURES

Location/Qualifiers

1..195

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/map="6p21.3"

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/protein_id="CAA58698.1"

/db_xref="GI:663003"

/db_xref="SPTREMBL:O19688"

/translation="IEQEGPEYWDRETQICKAKAQTDRDLRLRYNOSEA"

BASE COUNT 45 a 55 c 73 g 22 t

ORIGIN

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Quality: 44.00 Length: 10

Ratio: 4.889 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-14 x HSLAB27V2 ..

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147 CGAGAGGACCTGCGACCCCTGCTCGCTAC 176

seq_name: gb_pr3:HSAB27HLA1

seq_documentation_block:

LOCUS HSAB27HLA1 270 bp DNA PRI 08-JUL-1998
DEFINITION Homo sapiens MHC class I antigen HLA-B gene (HLA-B*27 variant allele) exon 2.

ACCESSION AF072763

VERSION AF072763.1 GI:3293562

KEYWORDS 1 of 2

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Homnidae; Homo.

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REFERENCE 1 (bases 1 to 270)
AUTHORS Steiner,N.K., Hurley,C.K. and Koester,R.P.
TITLE Novel-HLA-B allele
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 270)
AUTHORS Steiner,N.K., Hurley,C.K. and Koester,R.P.
TITLE Direct Submission
JOURNAL Submitted (21-JUN-1998) Microbiology and Immunology, Georgetown
University Medical Center, 3970 Reservoir Road NW, Washington, DC
20007, USA
FEATURES
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    exon            1..270
    /gene="HLA-B"
    /number=2
BASE COUNT      54 a 87 c 93 g 36 t
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    Ratio: 4.889        Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 90.000
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|||||
222 CGAGAGACCTCGGACCTGCTCCGCTAC 251
seq_name: gb_pr3:HSB1524V1
seq_documentation_block:
LOCUS HSB1524V1 270 bp DNA PRI 22-AUG-1998
DEFINITION Homo sapiens isolate GN00211 MHC class I antigen HLA-B gene (B*1543
allele), exon 2.
ACCESSION AF054011
VERSION AF054011.1 GI:2984766
KEYWORDS
SEGMENT
SOURCE
    1 of 2
    human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 270)
AUTHORS Kosman,C.A. and Hurley,C.K.
TITLE Novel HLA Class I B locus alleles
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 270)
AUTHORS Kosman,C.A. and Hurley,C.K.
TITLE Direct Submission
JOURNAL Submitted (18-MAR-1998) Microbiology & Immunology, Georgetown
University, 3970 Reservoir Rd. NW, Washington, DC 20007, USA
FEATURES
    source          Location/Qualifiers
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    /isolate="GN00211"
    /db_xref="taxon:9606"
    exon            1..270
    /gene="HLA-B"
    /number=2
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    Quality: 44.00      Length: 10
    Ratio: 4.889        Gaps: 0

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Percent Similarity: 90.000      Percent Identity: 90.000
alignment_block:
US-08-653-294-14 x HSB1524V1 ..
Align seg 1/1 to: HSB1524V1 from: 1 to: 270
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222 CGAGAGACCTCGGACCTGCTCCGCTAC 251
seq_name: gb_pr3:HSMHCHLBC1
seq_documentation_block:
LOCUS HSMHCHLBC1 270 bp DNA PRI 17-SEP-1998
DEFINITION Homo sapiens MHC class I antigen HLA-B gene (HLA-B*5303 allele),
exon 2.
ACCESSION AF071769
VERSION AF071769.1 GI:3243269
KEYWORDS
SEGMENT
SOURCE
    1 of 2
    human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 270)
AUTHORS Kosman,C.A. and Hurley,C.K.
TITLE Novel HLA Class I B Locus Alleles
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 270)
AUTHORS Kosman,C.A. and Hurley,C.K.
TITLE Direct Submission
JOURNAL Submitted (12-JUN-1998) Microbiology & Immunology, Georgetown
University, 3970 Reservoir Rd. N.W., Washington, DC 20007, USA
FEATURES
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    /isolate="GN00231"
    /db_xref="taxon:9606"
    exon            1..270
    /gene="HLA-B"
    /number=2
BASE COUNT      59 a 90 c 84 g 37 t
ORIGIN
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    Ratio: 4.889        Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 90.000
alignment_block:
US-08-653-294-14 x HSMHCHLBC1 ..
Align seg 1/1 to: HSMHCHLBC1 from: 1 to: 270
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222 CGAGAGACCTCGGACCTGCTCCGCTAC 251
seq_name: gb_pr4:HS27052B1
seq_documentation_block:
LOCUS HS27052B1 270 bp DNA PRI 21-JUN-1999
DEFINITION Homo sapiens MHC class I antigen HLA-B gene, HLA-B*2716 allele,
exon 2.
ACCESSION AF102563
VERSION AF102563.1 GI:4704574
KEYWORDS
SEGMENT
SOURCE
    1 of 2
    human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

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Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 270)
 AUTHORS Kosman,C.A. and Hurley,C.K.
 TITLE Novel Class I HLA-B Alleles
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 270)
 AUTHORS Kosman,C.A. and Hurley,C.K.
 TITLE Direct Submission
 JOURNAL Submitted (28-OCT-1998) Microbiology and Immunology, Georgetown University, 3970 Reservoir Rd. N.W., Washington, DC 20007, USA

FEATURES
 source Location/Qualifiers

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 /gene="HLA-B"
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 56 a 88 c 90 g 36 t
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 ORIGIN

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 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000

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Align seg 1/1 to: HS27052B1 from: 1 to: 270

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 222 CGAGAGGACCTGCGGACCTGCTCGCTAC 251

seq_name: gb_pr1:HS27052B1

seq_documentation_block:
 LOCUS HSLAWMBY2 270 bp DNA PRI 06-APR-1999
 DEFINITION Homo sapiens MHC class I antigen HLA-B gene (HLA-B*27 allele), exon

ACCESSION AF110257
 VERSION AF110257.1 GI:4566542

KEYWORDS
 SEGMENT 2 of 3
 SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 270)
 AUTHORS Wu,J., Bassinger,S., Montoya,G.D., Yee,J., Griffith,B.B.,
 Kearns,J., McKeen,M., Birkos,S., Kamoun,M. and Williams,T.M.

TITLE Identification of new HLA-B alleles in potential bone marrow donors
 JOURNAL Unpublished

REFERENCE 2 (bases 1 to 270)
 AUTHORS Wu,J., Bassinger,S., Montoya,G.D., Yee,J., Griffith,B.B.,
 Kearns,J., McKeen,M., Birkos,S., Kamoun,M. and Williams,T.M.
 TITLE Direct Submission
 JOURNAL Submitted (30-NOV-1998) Pathology, Univ. New Mexico, 915 Camino de
 Salud, NE, Albuquerque, NM 87131, USA

FEATURES
 source Location/Qualifiers

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 /db_xref="taxon:9606"
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 /gene="HLA-B"
 /number=2
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 ORIGIN

alignment_scores:

Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000

alignment_block:

US-08-653-294-14 x HSHLAWMBY2 ..
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 222 CGAGAGGACCTGCGGACCTGCTCGCTAC 251

seq_name: gb_pr2:HUMB27052G

seq_documentation_block:
 LOCUS HUMB27052G 546 bp DNA PRI 27-FEB-1996
 DEFINITION Homo sapiens MHC class I HLA-B*27052 gene, exons.

ACCESSION L76095

VERSION L76095.1 GI:1203957

KEYWORDS cell surface antigen; cell surface glycoprotein; class I gene;
 integral membrane protein; major histocompatibility complex.

SOURCE Homo sapiens DNA.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 546)

AUTHORS Marcos,C.Y., Fernandez-Vina,M.A., Lazaro,A.M. and Stastny,P.

TITLE Novel HLA-B Alleles

JOURNAL Unpublished (1996)

FEATURES Location/Qualifiers

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gene

1..270
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exon

271..546
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exon

allele

454
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gene

454
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BASE COUNT 109 a 172 c 195 g 70 t

ORIGIN

alignment_scores:

Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000

alignment_block:

US-08-653-294-14 x HUMB27052G ..
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 222 CGAGAGGACCTGCGGACCTGCTCGCTAC 251

seq_name: gb_pr1:HUMMBW2

seq_documentation_block:
 LOCUS HUMMBW2 822 bp DNA PRI 07-JAN-1995
 DEFINITION Human MHC class I HLA-B27 M+ gene, exons 2-4 (introns unsequenced).

ACCESSION M14013

VERSION M14013.1 GI:187743

KEYWORDS

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SOURCE      Human DNA.
ORGANISM    Homo sapiens
REFERENCE    Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
            Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS      1 (bases 1 to 822)
            Coppin,H.L. and McDevitt,H.O.
TITLE        Absence of polymorphism between HLA-B*27 genomic exon sequences
            isolated from normal donors and ankylosing spondylitis patients
JOURNAL      J. Immunol. 137 (7), 2168-2172 (1986)
MEDLINE      87009855
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  CDS
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    /note="HLA-B*27 M2+"
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    /db_xref="GDB:G00-120-048"
    /protein_id="AAA59643.1"
    /db_xref="GI:187744"
    /translation="GSHSMRYFHTSVSRGGERPRITVGVVDTLFLVRFDSDAASPR
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    VPGRLRGYHQDAYDGKDYIALNEDLSWTAADTAQITQKWEAAVAEQLRVYL
    EGECVEMLRYLENGKETLQADPPKTHVTHPTSDHEATLRCWALGFYPAEITLTWQ
    RDGEDOTDELVELTRPADRTFOKAAVVVPSGEQRYTCHVQHEGLPKPLTLRW"
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ORIGIN
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  Ratio: 4.889 Gaps: 0
  Percent Similarity: 90.000 Percent Identity: 90.000
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  223 CGAGAGGACCTCGGACCTGCTCCGCTAC 252
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seq_documentation_block:
LOCUS      HSU11267 968 bp mRNA PRI 21-JUL-1994
DEFINITION Human HLA-B*37 (B-3701) mRNA, partial cds.
ACCESSION  U11267
VERSION     U11267.1 GI:511785
KEYWORDS
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (sites)
AUTHORS      Hurley,C.K., Bel,M., Rodriguez,S. and Johnson,A.
TITLE        HLA-B*71
JOURNAL      Unpublished
REFERENCE    2 (bases 1 to 968)
AUTHORS      Hurley,C.K.
TITLE        Direct Submission
JOURNAL      Submitted (23-JUN-1994) Carolyn K. Hurley, Microbiology, Georgetown
            University School of Medicine, 3900 Reservoir Road, N.W.,
            Washington, D.C. 20007 USA
FEATURES
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DTAAQITQKWEAAVAEQLRVYLEGTCVEMLRYLENGKETLQADPPKTHVTHPT
SDHEATLRCWALGFYPAEITLTWQDGEDOTDELVELTRPADRTFOKAAVVVPSG
EEQRYTCHVQHEGLPKPLTLRWEPSSQSTIPVIGVAGLAVVV"
BASE COUNT  196 a 310 c 314 g 148 t
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  Percent Similarity: 90.000 Percent Identity: 90.000
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  Align seg 1/1 to: HSU11267 from: 1 to: 968
  1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
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  295 CGAGAGGACCTCGGACCTGCTCCGCTAC 324
seq_name: gb_pr1:HUMHMC
seq_documentation_block:
LOCUS      HUMHMC 1017 bp mRNA PRI 07-JAN-1995
DEFINITION Human MHC class I HLA-B*27-HS mRNA, 3' end.
ACCESSION  M62852
VERSION     M62852.1 GI:187760
KEYWORDS    cell surface antigen; class I gene; integral membrane protein;
            major histocompatibility complex.
SOURCE      Homo sapiens cDNA to mRNA.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE    1 (bases 1 to 1017)
AUTHORS      Choo,S.Y., Fan,L.A. and Hansen,J.A.
TITLE        A novel HLA-B*27 allele maps B27 allospecificity to the region
            around position 70 in the alpha 1 domain
JOURNAL      J. Immunol. 147 (1), 174-180 (1991)
MEDLINE     91268545
FEATURES
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    /product="MHC HLA-B*27-HS"
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    /db_xref="GI:187761"
    /translation="GSHSMRYFHTSVSRGGERPRITVGVVDTLFLVRFDSDAASPR

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EEPRAPWTEQSGPEVWDRETOICKAKAQTDRDLRLTLRYNQSGSHTLQSMYGC
VGPGRLLRGHNQAYDKDYIALNEDLRSWTAADTAQITQRKWEARVAEQRLRAYL
EGECVEMRLYLENGKETLQADPKTHVTHHPISDHEATLRCWALGFYPAEITLWQ
RGDEOTODTVELTRPADRTFOKWAAYVVPSEGEQRYTCHVOHEGLPKPLTLRWEP
SSQSTPIVGIVAGLAVLVVIGAVVAVMCRKSSGKGSYSQAACSDSAQGSDDV
SLTA"

BASE COUNT 207 a 308 c 343 g 159 t
ORIGIN

alignment_scores:
Quality: 44.00 Length: 10
Ratio: 4.889 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-14 x HUMHC ..
Align seg 1/1 to: HUMHC from: 1 to: 1017

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
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223 CGAGAGGACCTGCGACCCCTGCTCCGCTAC 252

seq_name: gb_pat:A28264

seq_documentation_block:
LOCUS A28264 1026 bp DNA PAT 24-MAY-1995
DEFINITION H.sapiens mRNA for HLA-B 27 from patent EP0226069.
ACCESSION A28264

VERSION A28264.1 GI:905320

KEYWORDS

SOURCE

human.

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;
Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 1026)

AUTHORS Szoets,H., Weiss,E., Doerner,C., Lang,M., Meo,T. and
Rietmueller,G.

TITLE HLA-B 27, DNA coding therefor and its utilization

JOURNAL Patent: EP 0226069-A 1 24-JUN-1987;

KEYWORDS Rietmueller, Gert, Prof. Dr

FEATURES Location/Qualifiers

source

1..1026

/organism="Homo sapiens"

/db_xref="taxon:9606"

BASE COUNT 213 a 307 c 344 g 162 t

ORIGIN

alignment_scores:
Quality: 44.00 Length: 10
Ratio: 4.889 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-14 x A28264 ..
Align seg 1/1 to: A28264 from: 1 to: 1026

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||
223 CGAGAGGACCTGCGACCCCTGCTCCGCTAC 252

seq_name: gb_pat:E01342

seq_documentation_block:
LOCUS E01342 1026 bp RNA PAT 29-SEP-1997
DEFINITION CDNA encoding C-terminal Fragment of HLA-B27.

ACCESSION E01342

VERSION E01342.1 GI:2169599

KEYWORDS JP 1987228281-A/2.

SOURCE Homo sapiens.

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Vertebrata; Mammalia; Eutheria;

Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 1026)

AUTHORS Hannecore,S., Eriabeeto,W., Kurisuta,D., Maagotsuto,R., Tomaso,M.

and Geruto,R.

TITLE HLA-B27, DNA ENCODING THE SAME AND ITS USE

JOURNAL Patent: JP 1987228281-A 2 07-OCT-1987;

BEHRINGWERKE AG

COMMENT

OS Human

PN JP 1987228281-A/2

PD 07-OCT-1987

PF 28-NOV-1986 JP 1986284078

PR 28-NOV-1985 DE 85 3542024, 21-DEC-1985 DE 85 3545576 PI

MAAGOTSUTO SUTSUETSU, ERIZABEETO WAISU, KURISUTA DERUNAA, PI

MAAGOTSUTO RANGU, TOMASO MEO, GERUTO RIITOMIYURAA PC

C12N15/00,C07H21/04,C12P21/00,C12Q1/68,G01N33/577//A61K39/00, PC

C07K13/00,

PC C07K15/06,(C12P21/00,C12R1:91);

CC strandedness: Double;

CC topology: Linear;

CC hypothetical: No;

CC anti-sense: No;

CC fragment_type: C-Terminal Fragment;

CC *source: cell_type=leukocyte;

CC Location/Qualifiers

Key

CD5

1..1026

/product='C-terminal fragment(exon 2 to 7)of

HLA-B27'.

FEATURES

source

1..1026

/organism="Homo sapiens"

/db_xref="taxon:9606"

BASE COUNT 213 a 307 c 344 g 162 t

ORIGIN

alignment_scores:

Quality: 44.00 Length: 10

Ratio: 4.889 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-14 x E01342 ..

Align seg 1/1 to: E01342 from: 1 to: 1026

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10

|||||

223 CGAGAGGACCTGCGACCCCTGCTCCGCTAC 252

seq_name: gb_pri:HUMMHZUNIA

seq_documentation_block:

LOCUS HUMMHZUNIA 1084 bp mRNA PRI 27-SEP-1993

DEFINITION Human MHC class I (HLA-B 27052) mRNA fragment.

ACCESSION L20086

VERSION L20086.1 GI:307282

KEYWORDS class I gene; lymphocyte antigen; major histocompatibility complex.

SOURCE Homo sapiens (strain South American Amerindian) cDNA to mRNA.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 1084)

AUTHORS Watkins,D.I., McAdam,S.N., Liu,X., Strang,C.R., Milford,E.L.,

Levine,C.G., Garber,T.L., Dogon,A.L., Lord,C.I., Ghim,S.H.,

Troup,G.M., Hughes,A.L. and Letwin,N.L.

New recombinant HLA-B alleles in a tribe of South American

Amerindians indicate rapid evolution of MHC class I loci

Nature 357, 329-333 (1992)

FEATURES

Location/Qualifiers

92269956

24

JOURNAL Unpublished
REFERENCE 2 (bases 1 to 1089)
AUTHORS Seurynck,K.L. and Baxter-Lowe,L.A.
TITLE Direct Submission
JOURNAL Submitted (22-SEP-1997) Molecular Genetics, Richland Memorial
Hospital, 7 Richland Medical Park, Columbia, SC 29203, USA
FEATURES
source
1..1089
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="6"
/cell_type="EBV transformed B-lymphocytes"
1..1089
/gene="HLA-B"
/allele="B27052"
1..1089
/gene="HLA-B"
/codon_start=1
/product="MHC class I antigen"
/protein_id="AAC42275.1"
/db_xref="GI:3643697"
/translation="MRVTEPRTLILLGVALTETWAGSHSMRYFHTSVSRPGRGE
RFITVGVDDTLFVRESDAASPREEPAPWIEQGEYWDRETOICKAKAQTDRDL
RTLLRYNQSEAGSHTLQNMVGCYDGPGRLLRGYHQDAYDGKDYIALNEDLSWTAA
DTAAQITQRKWEAKRAYEQLRAYLEGECEWLRRLRYLENGKETLQADPPKTHVTHPI
SDHEATLFCWALGFYPAEITLTWQDGEDQDTVELVETRPAGDRTFOKAAVVPVSG
EEQRYTCHVQHEGLPKPLTLRWEPSSQSTVPIVGIVAGLAVLVAVVIGAVVAVMCR
KSSGGKGSYSOAACSDSAQGSVSLTA"
BASE COUNT 217 a 332 C 368 g 172 t
ORIGIN

alignment_scores:
Quality: 44.00 Length: 10
Ratio: 4.889 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-14 x AF026218 ..
Align seg 1/1 to: AF026218 from: 1 to: 1089
1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||
295 CGAGAGGACCTCGGACCTGCTCCGCTAC 324

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Sequence	Strd	Orig	ZScore	EScore	Len	Documentation
N_Genes36.N70935	+	44.00	144.99	1.78	1026	Sequence encoding the human h
N_Genes36.N70225	+	44.00	133.50	7.78	3874	Sequence of genomic DNA encodi
N_Genes36.N71639	+	44.00	128.95	13.94	6553	HLA-B*27 consensus sequence. De
N_Genes36.N702167	+	39.00	138.25	4.23	270	HLA-B*52 exon 2 alpha-1 domain
N_Genes36.N701834	+	39.00	126.20	19.83	1086	Sequence encoding HLA-B*51 anti
N_Genes36.N701822	+	39.00	126.20	19.83	1086	Sequence encoding HLA-B*52 anti
N_Genes36.N705693	+	39.00	126.18	19.89	1089	HLA-B*51 gene for production of
N_Genes36.N7005701	+	39.00	126.18	19.89	1089	HLA-B*52 gene for production of
N_Genes36.N7012114	+	39.00	126.18	19.89	1089	HLA-B*53 exon. HLA-B*53 gene,
N_Genes36.N7078405	+	34.00	122.00	34.00	213	Human genome fragment. (Prefer
N_Genes36.N7012083	+	34.00	122.96	38.82	240	H.influenzae strain Minna (OMP
N_Genes36.N751732	+	34.00	108.80	184.62	978	H. influenzae a human secreted p
N_Genes36.N728520	+	34.00	104.55	318.39	1598	H. influenzae detection probe
N_Genes36.N74565	+	34.00	99.45	612.47	2881	Staphylococcus aureus contig S
N_Genes36.N767406	+	34.00	98.76	669.83	3123	Neural alpha-catenin protein c
N_Genes36.N765380	+	34.00	98.42	699.42	3247	Tomato ringspot virus peach is
N_Genes36.N719871	+	34.00	93.57	1.3e+03	5688	Rattus norvegicus cdo tumour s
N_Genes36.N719008	+	34.00	93.57	1.3e+03	5688	Rattus norvegicus cdo tumour s
N_Genes36.N719007	+	34.00	93.06	1.4e+03	6030	Rattus norvegicus Class II tum
N_Genes36.N719870	+	34.00	93.05	1.4e+03	6039	Rattus norvegicus cdo tumour s
N_Genes36.N731170	+	34.00	88.22	2.6e+03	10555	! Enterococcus faecalis genome
N_Genes36.N762176	+	34.00	67.38	3.6e+04	117213	! HSV-2 strain SB5 Contig ID 1
N_Genes36.N77006	+	33.00	119.37	47.62	189	Human genome fragment. New nu
N_Genes36.N707809	+	33.00	98.04	734.49	2223	Aspartokinase II gene. DNA enc
N_Genes36.N727730	+	33.00	97.89	748.08	2260	Insulin-stimulated protein kin
N_Genes36.N727731	+	33.00	97.89	748.08	2260	Mutant insulin-stimulated prot
N_Genes36.N7042561	+	33.00	95.96	1.0e+03	2960	Histamine H1 receptor coding s
N_Genes36.N700477	+	33.00	89.84	2.1e+03	5733	Arabidopsis thaliana clavata
N_Genes36.N720262	-	33.00	85.60	3.6e+03	9359	! Borrelia burgdorferi polyucle
N_Genes36.N715949	+	33.00	82.37	5.5e+03	13585	! Tumour rejection antigen pred
N_Genes36.N708946	+	32.00	125.78	20.91	59	U7.6 L3' PCR primer for U7.6 var
N_Genes36.N712619	+	32.00	123.15	29.32	80	! 2C11 scrv VL PCR primer 6. Singl
N_Genes36.N709941	+	32.00	122.83	30.55	83	! VK3'AL2 PCR primer for U7.6 var
N_Genes36.N762944	+	32.00	106.72	241.19	534	Sequence of carcinoembryonic an
N_Genes36.N750356	+	32.00	102.71	403.52	849	Sequence encoding fused antibod
N_Genes36.N713670	+	32.00	100.73	520.05	1067	! Enterococcus faecalis genome c
N_Genes36.N701865	+	32.00	100.36	545.53	1114	! Fc(epsilon) CH2'-CH4 coding se
N_Genes36.N721328	+	32.00	99.13	638.68	1284	Human C epsilon exon. New immu
N_Genes36.N7087474	+	32.00	98.97	651.95	1308	Human IgE Fc chain (amino acid
N_Genes36.N7091170	+	32.00	98.97	651.95	1308	Human IgE Fc chain (amino acid
N_Genes36.N705170	+	32.00	97.87	750.57	1485	Sequence encoding human immu
N_Genes36.N704062	+	32.00	97.87	750.57	1485	Sequence encoding human immu

```

FT intron 3009..3041
FT /*tag= f
FT intron 3148..3191
FT /*tag= g
PN EP-226069-A.
PD 24-JUN-1987.
PF 21-NOV-1986; 116139.
PR 01-JAN-1985; DE-542024.
PR 21-DEC-1985; DE-545576.
PR (BEHW ) BEHRINGER AG.
PI Szöcs H, Weiss E, Dorner C, Lang M, Meo T, Riethmüller G;
DR WPI; 87-171469/25.
DR P-PSDB; P70155.
DR DNA coding for human histocompatibility antigen HLA-B 27 - useful
PT for diagnosis and antigen and antibody prodn.
PS Claim 1; p6; 13pp; German.
CC The DNA may be used to detect the HLA-B 27 gene (opt. mutated) in
CC human genetic material. The HLA-B 27 may be used to detect anti-HLA-
CC B 27 antibodies in human serum. The antibodies may be used to
CC determine HLA-B 27 levels in human serum, e.g. for diagnosis of
CC rheumatic disorders, esp. ankylosing spondylitis.
SQ Sequence 3874 BP; 751 A; 1094 C; 1171 G; 858 T;

alignment_scores:
Quality: 44.00 Length: 10
Ratio: 4.889 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-14 x N70225 ..
Align seg 1/1 to: N70225 from: 1 to: 3874

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
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941 CGAGAGGACCTGCGGACCTGCTCGCTAC 970

seq_name: N_Geneseq_36:T61639

seq_documentation_block:
ID T61639 standard; DNA; 6553 BP.
AC T61639;
DT 05-JUN-1997 (first entry)
DE HLA B27 consensus sequence.
KW HLA B27; seronegative spondylarthropathy; ankylosing spondylitis;
KW Reiter's syndrome; arthritis; acute anterior uveitis; diagnosis;
KW ss; ds.
OS Homo sapiens.
FH Key Location/Qualifiers
FT mrna 3968..6653
FT /*tag= a
FT /note= "HLA-B27 3' flanking region, downstream of
FT 3' untranslated region"
FT mrna 4112..4556
FT /*tag= b
FT /note= "3' flanking region diagnostic for genetic
FT predisposition to SNSA"
FT mrna 4270..4556
FT /*tag= b
FT /note= "3' flanking region diagnostic for genetic
FT predisposition to SNSA"
FT misc_difference 4495
FT /*tag= d
FT /note= "absence of cytosine at this site is
FT indicative of a predisposition to SNSA"
PN W09709450-A1.
PD 13-MAR-1997.
PF 16-AUG-1996; U13256.
PR 01-SEP-1995; US-522942.
PR (CEDA-) CEDARS SINAI MEDICAL CENT.
PI Tyan DB;
DR WPI; 97-192924/17.

```

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PT Detecting pre-disposition to seronegative spondylarthropathies -
PT from the absence of a C residue at a specific position in the
PT 3'-flanking region of the HLA B27 allele
PS Claim 1; Page 52-56; 68pp; English
CC Genetic predisposition to seronegative spondylarthropathies (SNSA)
CC is detected by determining the absence of a cytosine nucleotide in
CC the 3' flanking region (see also T61647-48) of an HLA-B gene at a
CC position corresponding to nucleotide 4495 of the HLA-B27 consensus
CC sequence given in T61639. Probes and primers (see also T61640-46)
CC based on this region can be used in diagnostic assays to detect the
CC genetic predisposition to SNSA, and permit the distinction of B27+
CC individuals who are resistant to SNSA from B27+ normal individuals
CC who are susceptible (but as yet unaffected) to such diseases.
SQ Sequence 6553 BP; 1443 A; 1619 C; 2017 G; 1474 T;

alignment_scores:
Quality: 44.00 Length: 10
Ratio: 4.889 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-14 x T61639 ..
Align seg 1/1 to: T61639 from: 1 to: 6553

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||
1102 CGAGAGGACCTGCGGACCTGCTCGCTAC 1131

seq_name: N_Geneseq_36:Q29167

seq_documentation_block:
ID Q29167 standard; DNA; 270 BP.
AC Q29167;
DT 09-MAR-1993 (first entry)
DE HLA-Bw 52 exon 2 alpha-1 domain.
KW Human leukocyte antigen; transgenic; germ cells; somatic cells;
KW expression; ss.
PN J04091731-A.
PD 25-MAR-1992.
PF 03-AUG-1990; 207329.
PR 03-AUG-1990; JP-207329.
PR (OLYU ) OLYMPUS OPTICAL CO.
DR WPI; 92-342893/42.
PT Transgenic non-human mammalian HLA-Bw 52 gene - useful for
PT analysis of expression of gene structure, and prodn. of
PT mouse model of human disease
PS Disclosure; Fig 1; 8pp; Japanese.
CC The sequence shows the exon 2 alpha-1 domain of the human leukocyte
CC antigen-Bw 52 gene. The complete gene may be introduced into non-
CC human mammals, pref. rat or mouse, or their ancestors at the primary
CC developmental biological step via transplantation into the zygote or
CC embryo to generate transgenic non-human mammals incorporating the
CC HLA-Bw 52 gene in both their germ cells and somatic cells. Transgenic
CC non-human mammals contg. HLA-Bw 52 are useful for the analysis of
CC expression of the gene, its structure, and prodn. of mouse models of
CC human disease. See also Q29166-72.
SQ Sequence 270 BP; 59 A; 88 C; 86 G; 37 T;

alignment_scores:
Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:
US-08-653-294-14 x Q29167 ..
Align seg 1/1 to: Q29167 from: 1 to: 270

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||

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222 CGAGAGAACCTGGGATCGCGCTCCGCTAC 251

seq_name: N_Geneseq_36:Q01834

seq_documentation_block:

AC Q01834 standard; DNA; 1086 BP.
DT 19-MAR-1991 (first entry)
DE Sequence encoding HLA-B51 antigen.
KW Probe: HLA class I DNA; immunogen; ss.
OS Homo sapiens.
PN EP354580-A.
PD 14-FEB-1990.
PF 10-AUG-1989.
PR 11-AUG-1988; JP-200758.
PA (OLYU) Olympos Optical Co., Ltd.
PI Kano K, Takiguchi;
DR WPI: 90-046289/07.
PT New DNA for class I human leucocyte antigens and derived probes and
PT transformed cells, useful for DNA typing, as immunogens etc.
PS Claim 1; Page 11; 23pp; English.
CC The HLA class I DNA can be used as a source of probes for use in DNA
CC typing. Transformed cells, which are useful as immunogens, can be
CC obtained by introducing these DNAs into eucaryotic cells.
SQ Sequence 1086 BP; 224 A; 334 C; 356 G; 172 T;

alignment_scores:

Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-14 x Q01834 ..
Align seg 1/1 to: Q01834 from: 1 to: 1086
1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||:||||| |||||
294 CGAGAGAACCTGGGATCGCGCTCCGCTAC 323

seq_name: N_Geneseq_36:Q01822

seq_documentation_block:

ID Q01822 standard; DNA; 1086 BP.
AC Q01822;
DT 19-MAY-1991 (first entry)
DE Sequence encoding HLA-Bw52 antigen.
KW Probe: HLA class I DNA; immunogen; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT cds 1..1086
FT /*tag= a
FT EP-354580-A.
PD 14-FEB-1990.
PF 10-AUG-1989.
PR 11-AUG-1988; JP-200758.
PA (OLYU) Olympos Optical Co., Ltd.
PI Kano K, Takiguchi;
DR WPI: 90-046289/07.
DR P-PSDB; R03142.
PT New DNA for class I human leucocyte antigens and derived probes and
PT transformed cells, useful for DNA typing, as immunogens etc.
PS Claim 2; pp11-12; 23pp; English.
CC The HLA class I DNA can be used as a source of probes for use in DNA
CC typing. Transformed cells, which are useful as immunogens, can be
CC obtained by introducing these DNAs into eucaryotic cells.
SQ Sequence 1086 BP; 223 A; 335 C; 358 G; 170 T;

alignment_scores:

Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-14 x Q01822 ..
Align seg 1/1 to: Q01822 from: 1 to: 1086
1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||:||||| |||||
294 CGAGAGAACCTGGGATCGCGCTCCGCTAC 323

seq_name: N_Geneseq_36:Q05693

seq_documentation_block:

ID Q05693 standard; DNA; 1089 BP.
AC Q05693;
DT 03-JAN-1991 (first entry)
DE HLA-B51 gene for production of monoclonal antibodies.
KW Allotype specific monoclonal anti-HLA antibodies; hybridomas;
KW transgenic animals; HLA-B51 gene; ss.
FH Key Location/Qualifiers
FT exon 1..73
FT /*tag= a
FT /*number=1
FT /*tag= b
FT /*tag= b
FT /*number=2
FT /*note="alpha 1-domain"
FT 344..619
FT /*tag= c
FT /*number=3
FT /*note="alpha 2-domain"
FT 620..895
FT /*tag= d
FT /*number=4
FT /*note="alpha 3-domain"
FT 896..1012
FT /*tag= e
FT /*number=5
FT 1013..1042
FT /*tag= f
FT /*number=6
FT 1043..1089
FT /*tag= g
FT /*number=7
EP-383183-A.
PD 22-AUG-1990.
PF 07-FEB-1990; 102424.
PR 08-FEB-1989; JP-029313.
PA (OLYU) OLYMPUS OPTICAL KK.
PI Takiguchi M;
DR WPI: 90-255479/34.
PT Allotype specific monoclonal anti- HLA antibodies prodn. - using
PT hybridomas derived from transgenic animals carrying HLA gene and
PT immunised with HLA antigen of different allotype
PS Disclosure; Fig 1 A-G; 20pp; English.
CC The human HLA-B51 gene was injected into fertilised mouse eggs and
CC then these introduced into the uterus of a pseudo pregnant mouse.
CC The young were tested to ensure incorporation of the gene into the
CC chromosome, and one of them mated 3 times with a normal male to
CC produce 16 young, seven of which carried the HLA-B51 gene.
CC The transgenic offspring were immunised with HLA antigen.
CC The spleen lymphocytes were fused with myeloma cells. Hybridomas
CC producing antibodies were selected.
CC See also Q05701.
SQ Sequence 1089 BP; 224 A; 335 C; 357 G; 173 T;

alignment_scores:

Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

CC Preferred sequences exhibit no more than 90% homology to a human
CC sequence known per se.
SQ Sequence 213 BP; 76 A; 39 C; 41 G; 57 T;

alignment_scores:
Quality: 34.00 Length: 9
Ratio: 4.250 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 77.778

alignment_block:

US-08-653-294-14 x Q76405 ..

Align seg 1/1 to: Q76405 from: 1 to: 213

1 ArgGluAspLeuArgIleLeuLeuArg 9

|||||
160 AGGAGGAGTTTAAAGTAATGAAGA 186

seq_name: N_Geneseq_36:Q12083

seq_documentation_block:

ID Q12083 standard; DNA; 240 BP.

AC Q12083;

DT 26-JUL-1991 (first entry)

DE H. influenzae strain Minna (OMP subtype 1H)-cro-lacZ fusion

DE construct.

KW bacterial meningitis; vaccine; P1 gene; T-cell antigen; pRSM793;

KW outer membrane protein; ss.

OS Haemophilus influenzae.

FH Key Location/Qualifiers

FT cds

FT 1..234

FT /*tag= a

FT /product= partial cro-lacZ-strain Minna P1 OMP

FT fusion protein

PN W09106652-A.

PD 16-MAY-1991.

PF 31-OCT-1990; CA0374.

PR 31-OCT-1989; GB-024473.

PA (CONN-) CONNAUGHT LAB LTD.

PA (UNTW) WASHINGTON UNIV ST LOUIS.

PI Munson RS, Grass S, Chong P, Yang Y, Fahim R, McVerry P;

PI Klein M;

DR WPI; 91-164201/22.

PT Outer membrane protein of Haemophilus influenzae type B - used as

PT vaccine against infections, esp. in infants and for diagnosis

PS Disclosure: Fig 5; 33pp; English.

CC Plasmid pRSM793 contains only the 3' portion of the P1 gene. The

CC plasmid is derived from pRSM188 which contains the full-length P1

CC sequence. The cro-lacZ-ompP1 fusion protein produced from pRSM793

CC was recognised by rabbit and guinea pig P1-specific antisera in

CC immunoblot analyses.

CC See also R12446-R12455.

SQ Sequence 240 BP; 83 A; 44 C; 38 G; 75 T;

alignment_scores:

Quality: 34.00 Length: 10

Ratio: 3.778 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 60.000

alignment_block:

US-08-653-294-14 x Q12083/rev ..

Align seg 1/1 to reverse of: Q12083 from: 1 to: 240

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10

|||||

208 CGTAAGATTTCGTCGTGAGAGTAT 179

seq_name: N_Geneseq_36:X51732

seq_documentation_block:

ID X51732 standard; DNA; 978 BP.
AC X51732;
DT 17-JUN-1999 (first entry)
DE DNA encoding a human secreted protein.
KW Human secreted protein; cancer; immune disorder; infection;
KW inflammatory disorder; skin disorder; tumour; atherosclerosis;
KW restenosis; autoimmune disorder; Alzheimer's disease;
KW peripheral neuropathy; trauma; spinal cord injury; allergy;
KW hematopoietic disorder; skeletal disorder; neurological disorder;
KW arthritic disorder; asthma; immunodeficiency disease; AIDS;
KW transplant rejection; ss.

OS Homo sapiens.

PN W09911293-A1.

PD 11-MAR-1999.

PF 03-SEP-1998; U18360.

PR 12-SEP-1997; US-058974.

PR 05-SEP-1997; US-057626.

PR 05-SEP-1997; US-057663.

PR 05-SEP-1997; US-057669.

PR 12-SEP-1997; US-058666.

PR 12-SEP-1997; US-058667.

PR 12-SEP-1997; US-058973.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Brewer LA, Ebner R, Lafleur DW, Moore PA, Olsen HS,

PI Rosen GA, Ruben SM, Shi Y;

DR WPI; 99-204988/17.

DR P-PSDB; Y12945.

PT New isolated human genes and the secreted polypeptides they encode
PT - useful for diagnosis and treatment of e.g. neurological disorders,
PT tumours, immune disorders, inflammation or haematological disorders
PS Claim 1; Page 170; 215pp; English.

CC X51701-55 encode human secreted proteins. The polynucleotides and
CC their corresponding secreted polypeptides are useful for preventing,
CC treating or ameliorating medical conditions, e.g. by protein or gene
CC therapy. Pathological conditions can also be diagnosed by determining the
CC the amount of the new polypeptides in a sample or by determining the
CC presence of mutations in the new polynucleotides. Specific uses are
CC described for each polynucleotide, based on which tissues they are
CC most highly expressed in, and include developing products for the
CC diagnosis or treatment of cancer, immune disorders, infection,
CC inflammatory disorders, skin disorders, tumours, atherosclerosis,
CC restenosis, autoimmune disorders, Alzheimer's disease, peripheral
CC neuropathies, trauma, spinal cord injuries, allergy, hematopoietic
CC disorders, skeletal disorders, neurological disorders, arthritic
CC disorders, asthma, immunodeficiency diseases, AIDS and transplant
CC rejection. The polypeptides are also useful for identifying their
CC binding partners.

SQ Sequence 978 BP; 285 A; 256 C; 197 G; 240 T;

alignment_scores:

Quality: 34.00 Length: 10

Ratio: 3.778 Gaps: 0

Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-14 x X51732/rev ..

Align seg 1/1 to reverse of: X51732 from: 1 to: 978

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10

|||||

99 AGGAGGAGCTGAGATCCAACTGCGGTGG 70

seq_name: N_Geneseq_36:T28520

seq_documentation_block:

ID T28520 standard; DNA; 1598 BP.

AC T28520;

DT 02-APR-1997 (first entry)

DE H. influenzae detection probe #2.

KW Detection; probe; amplification primer; bacterial pathogen; pneumonia;

KW Escherichia coli; Klebsiella pneumoniae; Pseudomonas aeruginosa;

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FT FT      /*tag= a "these bases represent a line of missing text in
FT FT      /note=" the sequence listing in the specification. They
FT FT      are included to maintain the nucleotide numbering"
FT FT      given in the specification for this DNA sequence"
FT FT      2341. 2400
FT FT      /*tag= b
FT FT      /note=" these bases represent a line of missing text in
FT FT      the sequence listing in the specification. They
FT FT      are included to maintain the nucleotide numbering"
FT FT      given in the specification for this DNA sequence"
PN PN      EP-786519-A2.
PN PN      30-JUL-1997. PD PD
PN PN      07-JAN-1997. PD PD
PR PR      05-JAN-1996; US-009861.
PR PR      (HUMA-) HUMAN GENOME SCI INC.
PI PI      Barash SC, Choi GH, Dillion PJ, Fannon MR, Kunsch CA,
PI PI      Rosen CA;
PI PI      WPI: 97-374922/35.
PR PR
PT PT      Polynucleotide(s) and proteins derived from Staphylococcus aureus -
PT PT      stored on computer readable medium and used in the production of
PT PT      anti-S.aureus vaccines
PT PT
PS PS      Claim 1; Page 1047-1049; 3271pp; English.
PS PS
CC CC      This sequence represents one of 5191 Staphylococcus aureus DNA sequences
CC CC      of the invention. The DNA sequences are recorded on a computer readable
CC CC      medium, preferably selected from a floppy or hard disk, random access
CC CC      memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
CC CC      the S. aureus DNA sequences allows putative functions to be assigned so
CC CC      that protein-encoding or regulatory regions of commercial, therapeutic or
CC CC      industrial importance can be obtained. Specifically, sequences which are
CC CC      likely to encode antigens have been identified and these polypeptides can
CC CC      be used in a vaccine composition against S.aureus infection. The
CC CC      polypeptides can also be used in a kit for the immunodetection of
CC CC      S.aureus in a sample. S.aureus is implicated in numerous human diseases,
CC CC      including cellulitis, eyelid infections, food poisoning, osteomyelitis,
CC CC      skin and surgical wound infections, scalded skin syndrome, toxic shock
CC CC      syndrome, etc. Organisms transformed with the DNA sequences can be used
CC CC      for recombinant production of the polypeptides. The new DNA sequences
CC CC      (and their fragments) are useful as primers or probes for isolating
CC CC      homologues of any of the S.aureus DNA sequences contained on the
CC CC      computer readable medium.
SQ SQ      Sequence 2881 BP; 1049 A; 376 C; 505 G; 826 T

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alignment_scores:
  Quality: 34.00      Length: 9
  Ratio: 3.778       Gaps: 0
  Percent similarity: 100.000  Percent Identity: 66.667

alignment_block:
  US-08-653-294-14 x V74565/rev ..

Align seq 1/1 to reverse of: V74565 from: 1 to: 2881

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ATTN: T/I has
COPIES TO REVEAL OF
FROM: 10/20/2007

2 GluAspLeuArgIleLeuLeuArgTyr 10
:::|||||:::|||||:::|||||:::
48 GACGATTTTAAAAATCTGCTAAGATT 922

seq_name: N_Geneseq_36:Q67406

seq_documentation_block:
ID Q67406 standard; cDNA to mRNA; 3123 BP.

seq_documentation_block:
ID Q67406 standard; cDNA to mRNA; 3123 BP.
AC Q67406;

DE	19-APR-1995	(first entry)
DT	Neural alpha-catenin protein coding sequence.	
DE	Adhesion; neural alpha catenin; tumour; metastasis; disease;	
KW	autoimmune disease; infectious disease; dermal disease;	
KW	arteriosclerosis; ss.	
OS	Homo sapiens.	
OS	Location/Qualifiers	
Key	125..2845	
FT	/tag= a	
FT	/product= Neural alpha catenin.	

PN J06211898-A.
 PD 02-AUG-1994.
 PF 25-DEC-1992: 358026.
 PR 25-DEC-1992: JP-358026.
 PA (TAKI) TAKARA SHUZO CO LTD.
 DR WPI: 94-283359/35.
 DR P-PSDB; R58778.
 PT Neural alpha-catenin protein and DNA - useful in the control of
 PT cell adhesion, e.g., in treatment of tumour (metastasis) and
 PT autoimmune disease
 PS Disclosure: Page 10-14: 14pp; Japanese.
 CC The neural alpha catenin can be used for the treatment of diseases
 CC related to intercellular adhesion such as primary tumour, tumour
 CC metastasis, autoimmune diseases, infectious diseases, dermal
 CC diseases and arteriosclerosis.
 SQ Sequence 3123 BP; 920 A; 674 C; 827 G; 702 T;

alignment_scores:
 Quality: 34.00 Length: 9
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 66.667

alignment_block:

US-08-653-294-14 x Q67406/rev ..

Align seg 1/1 to reverse of: Q67406 from: 1 to: 3123

2 GluAspLeuArgIleLeuLeuArgTyr 10
 |||||
 1390 GAAGACCTGGCGGTACTCCTTCACTTC 1364

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OM of: US-08-653-294-14 to: EST:* out_format : pfs
 Date: Feb 8, 2000 4:02 AM
 About: Results were produced by the GenCore software, version 4.5,
 Copyright (c) 1993-2000 CompuGen Ltd.

Command line parameters:

-MODEL=framet_p2n_model -DEV=xlp
 -O=cpnl1/USPTO_spool/US08653294/runat_04022000_160700_15770/app_query.fasta.1
 -DB=EST -QFMT=fastap -SUFFIX=rst -GAPOP=12.000 -GAPEXT=4.000
 -MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000 -GAPOP=4.500
 -FCAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
 -FCAPEXT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DELOP=6.000
 -DELEXT=7.000 -START=1 -MATRIX=blosum62 -TRANS=human40.cdi
 -LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
 -OUTFMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=1000000 -USER=US08653294
 -NCPUP=6 -ICPU=3 -NO_XLPHY -WAIT -THREADS=1

Search information block:

Query: US-08-653-294-14
 Query length: 10
 Database: EST:*
 Database sequences: 4538634
 Database length: 1887831982
 Search time (sec): 8553.360000

score_list:

Sequence	Strd	Orig	ZScore	Escore	Len	Documentation
gb_est38:AL036690	+	44.00	165.79	1.81	171	AL036690 DKFZ564D2463.r1.564
gb_est30:AU056838	-	41.00	142.61	35.40	701	AU056838 Oryza sativa
gb_est21:AA989542	-	40.00	143.34	32.21	402	AA989542 sm64802.s1 Barstead SH
gb_gss13:AO440876	-	40.00	141.46	40.65	501	AQ440876 HS-5098.B2.B04.T7A RCP
gb_est21:AA975627	-	40.00	141.46	41.00	505	AA975627 og63805.s1 NCI_CGAP.k1
gb_est8:CO3945	+	39.00	144.03	29.51	232	CO3945 CO3945 Human heart cDNA
gb_est10:AA151891	+	39.00	143.25	32.61	255	AA151891 z001f06.r1 Stratagene
gb_est11:AA952680	+	39.00	142.81	34.50	269	AA952680 TENS1864.T1. cruzi epim
gb_est11:AA263158	+	39.00	142.39	36.41	283	AA263158 PMY0534 KGI-a Lambda Z
gb_est6:D82221	+	39.00	140.07	49.03	375	D82221 HUMHBC4626 Human pancrea
gb_gss13:AO44169	-	39.00	139.51	52.63	401	AQ44169 GSSTC0231 Trypanosoma
gb_est37:AI957215	+	39.00	139.05	55.83	424	AI957215 ul77a10.x1 Sugano mous
gb_est10:AA147151	+	39.00	136.46	77.89	581	AA147151 z032806.r1 Stratagene
gb_est26:AT359260	-	39.00	135.95	83.15	618	AT359260 GY27807.x1 NCI_CGAP.B
gb_gss13:AO449604	+	39.00	134.75	97.01	715	AQ449604 500002D08.x2 Cp10WAM3
gb_est31:AI698864	+	39.00	134.37	101.75	748	AI698864 C74h11.x1 NCI_CGAP.Pa
gb_gss8:AA039738	+	38.00	136.11	81.45	380	AA039738 CIT-HSP-2317E17.TF CIT
gb_est37:AI946856	-	38.00	135.28	90.55	420	AI946856 bs31h08.y1 Drosophila
gb_gss13:AO440598	-	38.00	135.09	92.83	430	AQ440598 HS-5089.B1.C10.SP6E RF
gb_est22:AT028215	-	38.00	135.01	93.74	434	AT028215 ov96c07.x1 Soares test
gb_gss11:AO301014	-	38.00	134.61	98.77	456	AQ301014 HS-3105.A2.E01.MR CIT
gb_est9:AA082472	-	38.00	134.46	100.61	464	AA082472 zn48a08.r1 Stratagene
gb_gss3:BB1151	+	38.00	134.18	104.28	480	B61151 T2005TF TAMU Arabidopsis
gb_est3:AA082478	-	38.00	132.72	125.75	573	AA082478 zn40507.r1 Stratagene
gb_est37:AI946939	-	38.00	132.47	129.94	591	AI946939 bs33c03.y1 Drosophila
gb_est24:AT239094	-	38.00	131.90	139.72	633	AT239094 GH15272.5prime GH Dros
gb_gss8:AO080015	+	37.00	133.78	109.80	316	AQ080015 CIT-HSP-2367F12.TR CIT
gb_est20:AA879637	+	37.00	133.01	121.22	347	AA879637 vx38b05.r1 Stratagene
gb_est10:AA178827	+	37.00	132.50	129.36	369	AA178827 mt68h02.r1 Soares mous
gb_est8:AA015279	-	37.00	131.80	141.62	402	AA015279 mh33d03.r1 Soares mous
gb_est32:AT748514	+	37.00	131.49	147.22	417	AT748514 sb54a12.y1 Gm-cl016 GL
gb_est28:AT508196	-	37.00	131.41	148.71	421	AT508196 mh33d03.y1 Soares mous
gb_est11:AA239196	-	37.00	131.14	153.94	435	AA239196 mx89c04.r1 Soares mous
gb_gss3:BA6971	+	37.00	130.96	157.69	445	BA6971 HS-1066.A2-E07-MR.abi CH
gb_gss13:AO435812	+	37.00	130.92	158.44	447	AQ435812 HS-5063.A1.B10.SP6E RF
gb_est16:AA596937	+	37.00	130.76	161.81	456	AA596937 vol14h04.r1 Barstead SH
gb_est16:AA592217	+	37.00	130.54	166.32	468	AA592217 vol1e03.r1 Barstead SH
gb_gss10:AO19154	-	37.00	129.98	178.75	501	AQ19154 RPCI11-45E5.TK RPCI1-11
gb_gss12:AO370195	-	37.00	129.75	184.03	515	AQ370195 HS-5045.B1.E03.T7 RPCI
gb_est22:AO014732	+	37.00	129.26	196.15	547	AO014732 AO014732 Mouse two-cel
gb_est25:AO045014	+	37.00	128.82	207.54	577	AO045014 AO045014 Mouse sixteen
gb_est39:AW119564	+	37.00	128.63	212.49	590	AW119564 sd48a12.y1 Gm-cl016 GL

gb_est22:AI055656 + 37.00 127.97 231.19 639 ! AI055656 coau0004K01 Cotton
 gb_est44:AW208428 + 37.00 127.29 252.28 694 ! AW208428 uo60c03.x1 NCI_CGAP
 gb_gss3:B20285 + 37.00 126.40 282.75 773 ! B20285 T2017-T7 TAMU Arabido

seq_name: gb_est38:AL036690

seq_documentation_block: 171 bp mRNA EST 27-SEP-1999
 LOCUS AL036690
 DEFINITION DKFZ564D2463.r1.564 (synonym: hfbr2) Homo sapiens CDNA clone
 DKFZ564D2463.5', mRNA sequence.

ACCESSION AL036690
 VERSION AL036690.3 GI:5927859
 KEYWORDS EST.
 SOURCE human.

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 171)
 AUTHORS Duesterhoeft,A., Lauber,J., Mewes,H.W., Gassenhuber,J. and
 Wiemann,S.

TITLE EST (Duesterhoeft, et al.)
 JOURNAL Unpublished (1999)
 COMMENT On Jul 7, 1999 this sequence version replaced gi:5866258.

Contact: Duesterhoeft A

MIPS

Am Klopferspitz 18a D-82152 Martinsried, Germany

This is the 5' sequence of the clone insert

Clone from S. Wiemann, Molecular Genome Analysis, German Cancer
 Research Center (DKFZ); Email s.wiemann@dkfz-heidelberg.de;
 sequenced by Qiagen within the CDNA sequencing consortium of the
 German Genome Project.

No sl sequence available.

This clone is available at the RZPD in Berlin.

Please contact the RZPD: Ressourcenzentrum, Heubnerweg 6, 14059

Berlin-Charlottenburg, GERMANY; Email: clone@rzpd.de.

FEATURES

Location/Qualifiers
 source
 1..171
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="DKFZ564D2463"
 /clone_lib="564 (synonym: hfbr2)"
 /tissue_type="brain"
 /dev_stage="fetal"
 /lab_host="xl-2blue"
 /notes="Vector: pAMP1; Site_1: NotI; Site_2: SalI"

BASE COUNT 36 a 53 c 60 g

ORIGIN

alignment_scores:

Quality: 44.00 Length: 10
 Ratio: 4.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-14 x AL036690 ..

Align seg 1/1 to: AL036690 from: 1 to: 171

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
 |||||
 62 CGAGAGGACCTGCGACCTGCTCGCTAC 91

seq_name: gb_est30:AU056838

seq_documentation_block: 701 bp mRNA EST 29-APR-1999
 LOCUS AU056838
 DEFINITION AU056838 Oryza sativa mature leaf Nipponbare Oryza sativa CDNA

clone S20919-1A, mRNA sequence.

ACCESSION AU056838

VERSION AU056838.1 GI:4715722

KEYWORDS EST.

SOURCE Oryza sativa.

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ORGANISM      Oryza sativa
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
Poaceae; Oryza.
1 (bases 1 to 701)
Yamamoto, K. and Sasaki, T.
Rice cDNA from mature leaf
Unpublished (1999)
On Jun 5, 1998 this sequence version replaced gi:3187083.
Contact: Takuji Sasaki
National Institute of Agrobiological Resources
Rice Genome Research Program
2-1-2 Kannondai, Tsukuba
Ibaraki,
Japan 305
Tel.: 0298-38-7441
Fax: 0298-38-7468
Email: tsasaki@agr.affrc.go.jp
PROJECT = "RGP".

FEATURES             source
    Location/Qualifiers
        1..701
        /organism="Oryza sativa"
        /strain="Nipponbare"
        /db_xref="taxon:4530"
        /clone="S20919_1A"
        /clone_lib="Oryza sativa mature leaf Nipponbare"
        /tissue_type="mature leaf"

BASE COUNT      145 a 169 c 230 g 151 t      6 others
ORIGIN

alignment_scores:
    Quality: 41.00      Length: 10
    Ratio: 4.556      Gaps: 0
    Percent Similarity: 90.000      Percent Identity: 90.000

alignment_block:
US-08-653-294-14 x AU056838/rev ..
Align seg 1/1 to reverse of: AU056838 from: 1 to: 701

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||
92 AGCGTGATCTCAGAAATCTCTTGAGATAC 63

seq_name: gb_est21:AA989542

seq_documentation_block:
LOCUS      AA989542      402 bp      mRNA      EST      02-JUN-1998
DEFINITION      am64d02.s1 Barstead spleen HPLRB2 Homo sapiens cDNA clone
                  IMAGE:1576803 3' similar to gb:L05093 60S RIBOSOMAL PROTEIN L18A
                  (HUMAN);, mRNA sequence.
ACCESSION      AA989542
VERSION        AA989542.1 GI:3174906
KEYWORDS       EST.
SOURCE         human.
ORGANISM       Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 402)
Hillier, L., Allen, M., Bowles, L., Dubuque, T., Geisler, G., Jost, S.,
Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M.,
Martin, J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F.,
Theising, B., White, Y., Wylie, T., Waterston, R. and Wilson, R.
WashU-NCI human EST Project
Unpublished (1997)
On Jan 19, 1998 this sequence version replaced gi:2153091.
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel.: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu

This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Trace considered overall poor quality
Seq primer: -40ml3 fwd. EX from Amersham
High quality sequence stop: 1.

FEATURES             source
    Location/Qualifiers
        1..402
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /clone="IMAGE:1576803"
        /clone_lib="Barstead spleen HPLRB2"
        /sex="male"
        /dev_stage="adult, 17 years"
        /lab_host="DH10B"
        /note="Organ: spleen; Vector: pT7T3D-Pac (Pharmacia) with
a modified polylinker; Site_1: EcoRI; Site_2: NotI; 1st
strand cDNA was primed with a Not I oligo(dT) primer [5'
TGTTACGAATCTGAAGTGGAGCGCCCTTTTCTTTTCTTTTCTTTTCTTTTCTTTT
3']; double-stranded cDNA was ligated to Eco RI adaptors
[AATTCGATCCTTG], digested with Not I and cloned into the
Not I and Eco RI sites of the modified pT7T3 vector.
Library constructed by Bob Barstead."

BASE COUNT      72 a 106 c 132 g 92 t
ORIGIN

alignment_scores:
    Quality: 40.00      Length: 9
    Ratio: 4.444      Gaps: 0
    Percent Similarity: 100.000      Percent Identity: 88.889

alignment_block:
US-08-653-294-14 x AA989542/rev ..
Align seg 1/1 to reverse of: AA989542 from: 1 to: 402

2 GluAspLeuArgIleLeuLeuArgTyr 10
|||||
353 GAAGAACTCCGATCTTCTGCGCTAT 327

seq_name: gb_gss13:AQ440876

seq_documentation_block:
LOCUS      AQ440876      501 bp      DNA      GSS      31-MAR-1999
DEFINITION      HS_5098_B2_B04_T7A RPCI-11 Human Male BAC Library Homo sapiens
                  genomic clone Plate=674 Col=8 Row=D, genomic survey sequence.
ACCESSION      AQ440876
VERSION        AQ440876.1 GI:4552215
KEYWORDS       GSS.
SOURCE         human.
ORGANISM       Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 501)
Mahairas, G.G., Wallace, J.C., Smith, K., Swartzell, S., Holzman, T., and
Keller, A., Shaker, R., Furlong, J., Young, J., Zhao, S., Adams, M.D. and
Hood, L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
99380589
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:

```

http://www.htsc.washington.edu
 Plate: 674 row: D column: 8
 Seq primer: T7
 Class: BAC ends
 High quality sequence stop: 501.

FEATURES

source
 1. .501
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="plate=674 Col=8 Row=D"
 /clone_lib="RPCr-11 Human Male BAC Library"
 /sex="male"
 /note="Vector: pBACE3.6; Genomic sequence of BAC ends"
 BASE COUNT 168 a 102 c 80 g 144 t 7 others
 ORIGIN

alignment_scores:
 Quality: 40.00 Length: 10
 Ratio: 4.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 60.000

alignment_block:

US-08-653-294-14 x A0440876/rev ..

Align seg 1/1 to reverse of: A0440876 from: 1 to: 501

1 ArgGluAspLeuArgIleLeuArgTyr 10
 |||||
 501 CGTGAAGACCTAAGAGTGTATACAGTTT 472

seq_name: gb_est21:AA975627

seq_documentation_block: 505 bp mRNA EST 22-MAY-1998
 LOCUS AA975627
 DEFINITION Oq63505.s1 NCI-CGAP_Kid6 Homo sapiens cDNA clone IMAGE:1590993 3' similar to gb:L05093 60S RIBOSOMAL PROTEIN L18A (HUMAN);, mRNA sequence.
 AA975627
 ACCESSION
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM

human.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 505)
 NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 Unpublished (1997)
 On May 8, 1995 this sequence version replaced gi:801263.
 Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550

FEATURES
 source
 Email: Robert.Strausberg@nih.gov
 Tissue Procurement: L. Jeffrey Medeiros, M.D., Michael R.
 Emmert-Buck, M.D., Ph.D.
 CDNA Library Preparation: Stratagene, Inc.
 CDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 www-bio.llnl.gov/bbrp/image/image.html

Seq primer: -40ml3 fwd. ET from Amersham
 High quality sequence stop: 255.

FEATURES

source
 1. .505
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:1590993"
 /clone_lib="NCI-CGAP_Kid6"
 /sex="mixed"
 /tissue_type="kidney tumor"

/lab_host="SOLR (kanamycin resistant)"
 /note="Organ: kidney; Vector: Bluescript SK-; Site: 1;
 EcoRI; Site 2: XhoI; Cloned unidirectionally. Primer:
 Oligo dt. Pooled kidney tumors. 5' adaptor sequence: 5'
 GAATTCGGCACGAG 3' 3' adaptor sequence: 5'
 CTCGAGCTTTTTCCTTTT 3' Average insert size: 1.0 kb."
 BASE COUNT 93 a 115 c 177 g 120 t
 ORIGIN

alignment_scores:
 Quality: 40.00 Length: 9
 Ratio: 4.444 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 88.889

alignment_block:

US-08-653-294-14 x AA975627/rev ..

Align seg 1/1 to reverse of: AA975627 from: 1 to: 505

2 GluAspLeuArgIleLeuArgTyr 10
 |||||
 314 GAAGAACTTCGATTCTGCTCGCTAT 288

seq_name: gb_est8:C03945

seq_documentation_block: 232 bp mRNA EST 30-JUL-1996
 LOCUS C03945
 DEFINITION C03945 Human heart cDNA (YNakamura) Homo sapiens cDNA clone
 3NHC2454, mRNA sequence.
 ACCESSION C03945
 VERSION C03945.1 GI:1467196
 KEYWORDS EST.
 SOURCE human.

ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 232)
 Tanaka,T., Ogiwara,A., Uchiyama,I., Takagi,T., Yazaki,Y. and
 Nakamura,Y.
 TITLE Construction of a normalized directionally cloned cDNA library from
 adult heart and analysis of 3040 clones by partial sequencing
 Genomics 35 (1), 231-235 (1996)
 96299762
 COMMENT On Oct 24, 1995 this sequence version replaced gi:1040105.
 Contact: Yusuke Nakamura
 Institute of Medical Science
 University of Tokyo
 4-6-1, Shirokanedai, Minato-ku, Tokyo 108, Japan
 Tel: 81-3-5449-5372
 Fax: 81-3-5449-5433
 Email: yusuke@ims.u-tokyo.ac.jp.

FEATURES
 source
 1. .232
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="3NHC2454"
 /clone_lib="Human heart cDNA (YNakamura)"
 /dev_stage="adult"
 /note="Organ: heart; normalized directionally cloned cDNA
 from adult heart"

BASE COUNT 55 a 77 c 68 g 32 t
 ORIGIN

alignment_scores:

Quality: 39.00 Length: 10
 Ratio: 4.333 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-14 x C03945 ..

Align seg 1/1 to: C03945 from: 1 to: 232

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||:|||||
40 CGAGAGAACCTGGGATCGGCTCCGCTAC 69

seq_name: gb_est10:AA151891

seq_documentation_block: 255 bp mRNA EST 10-DEC-1996
LOCUS AA151891
DEFINITION Z001f06.r1 Stragatene colon (#937204) Homo sapiens cDNA clone IMAGE:566435 5' similar to gb:M15497_cds1 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN, A-24(A-9) A*2401 (HUMAN);, mRNA sequence.
ACCESSION AA151891
VERSION AA151891.1 GI:1720754
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 255)
AUTHORS Hillier,L., Lennon,G., Becker,M., Bonaldo,M.F., Chiapelli,B., Chisoe,S., Dietrich,N., Dubouche,T., Favello,A., Gish,W., Hawkins,M., Hultman,M., Kucaba,T., Lacy,M., Le,M., Le,N., Marais,E., Moore,B., Morris,M., Parsons,J., Prange,C., Rifkin,L., Rohlfing,T., Schellenberg,K., Soares,M.B., Tan,F., Thierry-Mieg,J., Trevaskis,E., Underwood,K., Wohlmann,P., Waterston,R., Wilson,R. and Marra,M.
TITLE Generation and analysis of 280,000 human expressed sequence tags
JOURNAL Genome Res. 6 (9), 807-828 (1996)
MEDLINE 9704478
COMMENT On May 8, 1995 this sequence version replaced gi:800234.
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
This clone is available royalty-free through LNL; contact the IMAGE Consortium (info@image.lnl.gov) for further information.
Trace considered overall poor quality
Seq primer: -28M13 rev2 from Amersham
High quality sequence stop: 1.
Location/Qualifiers
1..255
/organism="Homo sapiens"
/db_xref="GDB:459088"
/db_xref="taxon:9606"
/clone="IMAGE:566435"
/lab_host="SOUR cells (kanamycin resistant)"
/note="organ: colon; Vector: pBluescript SK-; Site:1: EcoRI; Site:2: XhoI; Cloned unidirectionally. Primer: Oligo dt. T-84 colonic epithelial cell line. Average insert size: 1.0 kb; Uni-ZAP XR Vector; -5' adaptor sequence: 5' GAATTCGACGACGAG 3' -3' adaptor sequence: 5' CTCGAGTGTGTTTTTTTTT 3'

BASE COUNT 57 a 70 c 75 g 44 t 9 others
ORIGIN
alignment_scores:
Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000
alignment_block:
US-08-653-294-14 x AA151891 ..
Align seg 1/1 to: AA151891 from: 1 to: 255

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10

|||||:|||||
77 CGAGAGAACCTGGGATCGGCTCCGCTAC 106

seq_name: gb_est21:AA952680

seq_documentation_block: 269 bp mRNA EST 29-OCT-1998
LOCUS AA952680
DEFINITION TENS1864 T. cruzi epimastigote normalized cDNA Library Trypanosoma cruzi cDNA clone 1864 5', mRNA sequence.
ACCESSION AA952680
VERSION AA952680.1 GI:3115776
KEYWORDS EST.
SOURCE Trypanosoma cruzi.
ORGANISM Trypanosoma cruzi
REFERENCE 1 (bases 1 to 269)
AUTHORS Eukaryota; Euglenozoa; Kinetoplastida; Trypanosomatidae; Trypanosoma; Schizotrypanum.
TITLE Gene discovery through expressed sequence tag sequencing in trypanosoma cruzi
JOURNAL Infect. Immun. 66 (11), 5393-5398 (1998)
MEDLINE 99003155
COMMENT On Jan 17, 1998 this sequence version replaced gi:1900451.
Contact: Sanchez D.O.
Instituto de Investigaciones Biotecnologicas (Univ. Nac. de Gral San Martin)
Av. Gral Paz entre Albarcellos y Constituyentes, INTI edificio 24 CP(1650) San Martin, Prov. de BS AS Argentina
Tel: (54-1)752-9639 or (54-1)752-0021
Fax: (54-1)752-0021 or (54-1)752-9639
Email: dsanchez@inti.gov.ar
Seq primer: T7.
Location/Qualifiers
1..269
/organism="Trypanosoma cruzi"
/strain="Cl-Brenner"
/db_xref="taxon:5693"
/clone="1864"
/cell_type="epimastigote"
/note="cDNA library constructed with oligo dt primed epimastigote mRNA and cloned in pT7t318D phagemid with modified polylinker (PHARMACIA)"

BASE COUNT 47 a 60 c 89 g 69 t 4 others
ORIGIN

alignment_scores:
Quality: 39.00 Length: 10
Ratio: 3.900 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 60.000

alignment_block:
US-08-653-294-14 x AA952680 ..
Align seg 1/1 to: AA952680 from: 1 to: 269
1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
|||||:|||||
177 CGGAGAGAGTTGAGGCTATTATCCGCTAC 206

seq_name: gb_est11:AA263158

seq_documentation_block: 283 bp mRNA EST 02-JUL-1998
LOCUS AA263158
DEFINITION PM70534 KGI-a Lambda zap Express cDNA library Homo sapiens CDNA 5', mRNA sequence.
ACCESSION AA263158
VERSION AA263158.1 GI:1898964
KEYWORDS EST.
SOURCE human.
ORGANISM Homo sapiens

```

tel: 272-20-8856
fax: 272-20-8896
Email: jtakeda@sb.gunma-u.ac.jp.
FEATURES
    Location/Qualifiers
        source
            1..375
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone_lib="Human pancreatic islet"
                /note="Vector: Lambda ZAPII; Site_1: Eco RI; Site_2: Xho
I; mRNA was prepared from normal adult human islets. cDNA
was directionally synthesized from the xho I in the vector
to the EcoRI site. cDNA was size fractionated to remove
sequences <1000 bp in size."
BASE COUNT      75 a      124 c      118 g      55 t      3 others
ORIGIN

alignment_scores
    Quality:      39.00
    Ratio:        4.333
    Percent Similarity: 90.000
    Gaps:         0
    Percent Identity: 80.000
    Length:      10

```

```

alignment_block:
  US-08-653-294-14 x D82221  ..
  Align seg 1/1 to: D82221 from: 1 to: 375

  1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
    |||||:||||| |||||
  306 CGAGAGAACCTCGGATCGCGTCGNTAC 335

seq_name: gb_gss13:AQ444169

seq_documentation_block:
  LOCUS      AQ444169          401 bp      DNA
  DEFINITION  GSSTC0231 Trypanosoma cruzi random genomic library Trypanosoma
              cruzi genomic clone G10N2 5', genomic survey sequence.
  ACCESSION  AQ444169
  VERSION    AQ444169.1 GI:4555633
  KEYWORDS   GSS.
  SOURCE     Trypanosoma cruzi.
  ORGANISM   Trypanosoma cruzi

```

Trypanosoma; Schizotrypanum.
1 (bases 1 to 401)
AUTHORS Sanchez,D.O.
TITLE Trypanosoma cruzi random genomic sequences
JOURNAL Unpublished (1999)
COMMENT Contact: Sanchez D.O.
Instituto de Investigaciones Biotecnologicas (Univ. Nac. de Gral
San Martin)
Av. Gral Paz entre Albarellos Y Constituyentes, INTI edificio 24
CP(1650) San Martin, Prov. de BS AS. Argentina
Tel: (54-11)4752-0021
Fax: (54-11)4752-9639
Email: dsanchez@inti.gov.ar
Seq primer: T7
Class: shotgun.
Location/Qualifiers
1. .401
/organism="Trypanosoma cruzi"
/strain="Cl-Brenner"
/db_xref="taxon:5693"
/clone="G10N2"
/clone_lib="Trypanosoma cruzi random genomic library"
/cell_type="epimastigote"
/note="Vector: pBS(-) (PHARMACIA)"
111 a 129 c 93 g 65 t 3 others
BASE COUNT
ORIGIN
alignment_scores:
Quality: 39.00 Length: 10

alignment_block:

US-08-653-294-14 x AA147151

Align seg 1/1 to: AA147151 from: 1 to: 581

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
 |||||:|||||
 152 CGAGAGACCTGCGGATCGCTCGCTAC 181

seq_name: gb_est26:A1359260

seq_documentation_block:

LOCUS A1359260 618 bp mRNA 15-FEB-1999
 DEFINITION qv27b07.x1 NCI-CGAP_Brn23 Homo sapiens cDNA clone IMAGE:2013205 3'
 similar to gb:D332129 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
 AW-66(A-10) A*6601 ALPHA (HUMAN)), mRNA sequence.
 ACCESSION A1359260
 VERSION A1359260.1 GI:4110881
 KEYWORDS EST.
 SOURCE human.

ORGANISM

Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 618)
 NCI/NINDS-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.
 National Cancer Institute / National Institute of Neurological
 Disorders and Stroke, Brain Tumor Genome Anatomy Project
 (CGAP/BTGA), Tumor Gene Index
 Unpublished (1998)

JOURNAL

COMMENT

Contact: Robert Strausberg, Ph.D.
 Tel: (301) 496-1550
 Email: Robert.Strausberg@nih.gov
 Tissue Procurement: David N. Louis, M.D., Myrna R. Rosenfeld M.D.,
 Ph.D.
 cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
 Bonaldo, Ph.D.
 cDNA Library Arrayed by: Greg Lennon, Ph.D.
 DNA Sequencing by: Washington University Genome Sequencing Center
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 691 Std Error: 0.00

Seq primer: -400p from Gibco

High quality sequence stop: 458.

FEATURES

source

1..618
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="IMAGE:2013205"
 /clone_lib="NCI-CGAP_Brn23"
 /tissue_type="gliblastoma (pooled)"
 /lab_host="DH10B"

/note="Organ: brain; Vector: pT7T3D-Pac (Pharmacia) with a
 modified polylinker; Site_1: Not I; Site_2: Eco RI; 1st
 strand cDNA was primed with a Not I - oligo(dT) primer [5'
 TGTTACCAATCTGAAGTGGAGCGCGGCATATCTTTTTTTTTTTTTTTT
 T 3']; double-stranded cDNA was ligated to Eco RI
 adaptors (Pharmacia), digested with Not I and cloned into
 the Not I and Eco RI sites of the modified pT7T3 vector.
 Library is normalized, and was constructed by Bento
 Soares and M.Fatima Bonaldo."

BASE COUNT 128 a 171 c 182 g 137 t

ORIGIN

alignment_scores:

Quality: 39.00 Length: 10
 Ratio: 4.333 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-14 x A1359260/rev

Align seg 1/1 to reverse of: A1359260 from: 1 to: 618

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
 |||||:|||||
 319 CGAGAGACCTGCGGATCGCTCGCTAC 290

seq_name: gb_gss13:AQ449604

seq_documentation_block:

LOCUS AQ449604 715 bp DNA GSS 08-APR-1999
 DEFINITION 50000ZD08.x2 CpiOWAM13mpl18gDNA1 Cryptosporidium parvum genomic,
 genomic survey sequence.
 ACCESSION AQ449604
 VERSION AQ449604.1 GI:4578741
 KEYWORDS GSS.
 SOURCE Cryptosporidium parvum.
 ORGANISM Cryptosporidium parvum

Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;
 Cryptosporidiidae; Cryptosporidium.

REFERENCE

1 (bases 1 to 715)
 Hyman, R. W., Fung, E., Qin, F., Rowley, D. and Davis, R. W.
 Cryptosporidium parvum genome sequencing demonstration project
 Unpublished (1999)

JOURNAL

COMMENT

On Mar 23, 1999 this sequence version replaced gi:3325323.
 Contact: Hyman, R. W.
 Stanford DNA Sequencing and Technology Center
 Stanford University School of Medicine, Palo Alto
 855 California Avenue, Palo Alto, CA 94304, USA
 Tel: 650 812 1972
 Fax: 650 812 1975
 Email: hyman@sequence.stanford.edu
 For Annotation Data see <http://medsfgh.ucsf.edu/id/CpTags/home.html>
 Seq primer: M13(-21) forward
 Class: shotgun.

FEATURES

source

1..715
 /organism="Cryptosporidium parvum"
 /strain="IOWA"
 /db_xref="taxon:5807"
 /clone_lib="CpiOWAM13mpl18gDNA1"
 /lab_host="E. coli DH125"
 /note="Vector: M13mpl8; Site_1: Hind III; C. parvum (IOWA
 isolate) genomic DNA was hydrodynamically sheared to
 produce fragments having a tight size distribution between
 1.5 and 3 kb. Adaptors (pGTGACTCA/CRAACCACTGAGTp) were
 ligated to the randomly sheared gDNA fragments and
 PACTGTGTTG linkers were ligated to the Hind III-cleaved
 M13mpl8 vector. The adaptor-containing inserts were
 annealed and ligated to the vector and transformed into E.
 coli strain DH125. Recombinant phagemid clones from the
 first plating of the library were randomly selected for
 sequence analysis using the M13(-21) forward primer."

BASE COUNT 272 a 92 c 109 g 241 t

ORIGIN

alignment_scores:

Quality: 39.00 Length: 10
 Ratio: 4.333 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-14 x AQ449604

Align seg 1/1 to: AQ449604 from: 1 to: 715

1 ArgGluAspLeuArgIleLeuLeuArgTyr 10
 |||||:|||||
 191 AGAGAGATCTTAATTTGTTGACCGATAT 220

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GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2000, 01:29:38 ; Search time 122.56 Seconds
(without alignments)
1.933 Million cell updates/sec

Title: US-08-653-294-15
Perfect score: 49
Sequence: 1 YRLRLRLDER 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : A_Geneseq_36.*

Pred. NO. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	49	100.0	10	1 W47272	Immunomodulatory p
2	44	89.8	10	1 W47268	Immunomodulatory p
3	44	89.8	10	1 W47270	Immunomodulatory p
4	39	79.6	10	1 W47266	Immunomodulatory p
5	39	79.6	20	1 R92909	HLA-B2702 CTL modu
6	39	79.6	20	1 R92911	HLA-B2702 CTL modu
7	39	79.6	20	1 R92907	HLA-B2702 CTL modu
8	39	79.6	20	1 R95428	HLA-B2702 84-75-84
9	39	79.6	20	1 W33778	Immunomodulating d
10	39	79.6	20	1 W33779	Immunomodulating d
11	39	79.6	20	1 W33792	Peptide B2702.84-7
12	34	69.4	20	1 R92910	HLA-B2702 CTL modu
13	34	69.4	20	1 R92908	HLA-B2702 CTL modu
14	34	69.4	20	1 R95430	HLA-B2702 84-75T/7
15	34	69.4	20	1 W33791	Peptide B2702.84-7
16	34	69.4	20	1 W33793	Peptide B2702.84-7
17	33	67.3	60	1 W93813	Rice anthranilate
18	33	67.3	104	1 W93811	Rice anthranilate
19	33	67.3	577	1 W93815	Rice ASA first iso
20	33	67.3	577	1 W93810	Rice anthranilate
21	32	65.3	451	1 R27842	Human calcium chan
22	30	61.2	319	1 W12377	Regulatory factor
23	30	61.2	381	1 W98786	H. pylori GHPO 121
24	30	61.2	387	1 W62846	A partial gldAI pr
25	30	61.2	621	1 W62842	Helicobacter pylor
26	30	61.2	625	1 W89445	A gldAI protein se
27	30	61.2	775	1 W79193	Human Hrs-2 partia
28	30	61.2	851	1 R41333	113 KD ISGF-3alpha
29	30	61.2	851	1 R72077	Recognition factor
30	30	61.2	851	1 W03166	Human STAT2, New S
31	30	61.2	924	1 W79192	Rat Hrs-2 polypept
32	30	61.2	3079	1 R59926	GAP protein Ira2.
33	30	61.2	3224	1 W54235	Human Nup358 prote
34	29	59.2	6	1 W47264	Immunomodulatory p

Peptide #4 used in
H. pylori GHPO 54
S. aureus gldB pro
S. aureus gldB pro
Rat FRAG1 protein.
3-acylation enzyme
EHV-4 gc. Nucleic
Human TIE ligand N
Mouse Smad6 protei
Smad7 protein used
Bacillus species a

ALIGNMENTS

RESULT 1

W47272
ID W47272 standard; peptide; 10 AA.
AC W47272;
DE 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.
FH Key Location/Qualifiers
FT Misc_difference 1..10 "at least one of the amino acids is the
FT D-isomer

PN W09744052-A1.
PD 27-NOV-1997.
PF 23-APR-1997; U06705.
PR 22-MAY-1996; US-651650.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI; 98-018220/02.
PT Novel immunomodulatory peptide-type compound - useful for inhibiting
PT transplant rejection

PS Claim 10; Page 36; 41pp; English.
CC The present sequence is an immunomodulatory peptide, which
CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
CC in a pharmaceutical composition together with a subtherapeutic dose
CC of an immunosuppressant, to extend the period of acceptance of a
CC transplant from a major histocompatibility complex (MHC) unmatched
CC donor, i.e. to inhibit transplant rejection. It can also be used in
CC the treatment of autoimmune diseases.
CC Peptides using the D-form amino acids are more effective
CC Immunomodulators than their diastereomers or enantiomers.
SQ Sequence 10 AA;

Query Match 100.0%; Score 49; DB 1; Length 10;
Best Local Similarity 100.0%; Pred. No. 0.00025;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 YRLRLRLDER 10
| | | | | | | | | |
DB 1 YRLRLRLDER 10

RESULT 2

W47268
ID W47268 standard; peptide; 10 AA.
AC W47268;
DE 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.
FH Key Location/Qualifiers
FT Misc_difference 1..10

```

FT      /note= "at least one of the amino acids is the
FT      D-isomer
PN      WO9744052-A1.
PD      27-NOV-1997.
PF      23-APR-1997; U06705.
PR      22-MAY-1996; US-651650.
PA      (STRD ) UNIV LELAND STANFORD JUNIOR.
PI      Clayberger C, Krensky AM;
DR      WPI; 98-018220/02.
PT      Novel immunomodulatory peptide-type compound - useful for inhibiting
PT      transplant rejection
PS      Claim 10; Page 36; 41pp; English.
CC      The present sequence is an immunomodulatory peptide, which
CC      comprises a Class I HLA-B alpha-1 domain sequence. It can be used
CC      in a pharmaceutical composition together with a subtherapeutic dose
CC      of an immunosuppressant, to extend the period of acceptance of a
CC      transplant from a major histocompatibility complex (MHC) unmatched
CC      donor, i.e. to inhibit transplant rejection. It can also be used in
CC      the treatment of autoimmune diseases.
CC      Peptides using the D-form amino acids are more effective
CC      immunomodulators than their diastereomers or enantiomers.
SQ      Sequence 10 AA;

Query Match      89.8%; Score 44; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.0027;
Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      1 YRLRLRLDER 10
DB      1 YRLRLRLNER 10

RESULT      3
W47270
ID      W47270 standard; peptide; 10 AA.
AC      W47270;
DT      22-MAY-1998 (first entry)
DE      Immunomodulatory peptide.
KW      Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW      transplant rejection; treatment; autoimmune disease.
OS      Homo sapiens.
FH      Key
FT      Key
FT      Location/Qualifiers
FT      Misc_difference 1..10
FT      /note= "at least one of the amino acids is the
FT      D-isomer
PN      WO9744052-A1.
PD      27-NOV-1997.
PF      23-APR-1997; U06705.
PR      22-MAY-1996; US-651650.
PA      (STRD ) UNIV LELAND STANFORD JUNIOR.
PI      Clayberger C, Krensky AM;
DR      WPI; 98-018220/02.
PT      Novel immunomodulatory peptide-type compound - useful for inhibiting
PT      transplant rejection
PS      Claim 10; Page 36; 41pp; English.
CC      The present sequence is an immunomodulatory peptide, which
CC      comprises a Class I HLA-B alpha-1 domain sequence. It can be used
CC      in a pharmaceutical composition together with a subtherapeutic dose
CC      of an immunosuppressant, to extend the period of acceptance of a
CC      transplant from a major histocompatibility complex (MHC) unmatched
CC      donor, i.e. to inhibit transplant rejection. It can also be used in
CC      the treatment of autoimmune diseases.
CC      Peptides using the D-form amino acids are more effective
CC      immunomodulators than their diastereomers or enantiomers.
SQ      Sequence 10 AA;

Query Match      89.8%; Score 44; DB 1; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.0027;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY      1 YRLRLRLDER 10
DB      1 YRLRLRLNER 10

RESULT      4
W47266
ID      W47266 standard; peptide; 10 AA.
AC      W47266;
DT      22-MAY-1998 (first entry)
DE      Immunomodulatory peptide.
KW      Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW      transplant rejection; treatment; autoimmune disease.
OS      Homo sapiens.
FH      Key
FT      Key
FT      Location/Qualifiers
FT      Misc_difference 1..10
FT      /note= "at least one of the amino acids is the
FT      D-isomer
PN      WO9744052-A1.
PD      27-NOV-1997.
PF      23-APR-1997; U06705.
PR      22-MAY-1996; US-651650.
PA      (STRD ) UNIV LELAND STANFORD JUNIOR.
PI      Clayberger C, Krensky AM;
DR      WPI; 98-018220/02.
PT      Novel immunomodulatory peptide-type compound - useful for inhibiting
PT      transplant rejection
PS      Claim 10; Page 36; 41pp; English.
CC      The present sequence is an immunomodulatory peptide, which
CC      comprises a Class I HLA-B alpha-1 domain sequence. It can be used
CC      in a pharmaceutical composition together with a subtherapeutic dose
CC      of an immunosuppressant, to extend the period of acceptance of a
CC      transplant from a major histocompatibility complex (MHC) unmatched
CC      donor, i.e. to inhibit transplant rejection. It can also be used in
CC      the treatment of autoimmune diseases.
CC      Peptides using the D-form amino acids are more effective
CC      immunomodulators than their diastereomers or enantiomers.
SQ      Sequence 10 AA;

Query Match      79.6%; Score 39; DB 1; Length 10;
Best Local Similarity 80.0%; Pred. No. 0.029;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY      1 YRLRLRLDER 10
DB      1 YRLRLRLNER 10

RESULT      5
R92909
ID      R92909 standard; peptide; 20 AA.
AC      R92909;
DT      16-MAY-1996 (first entry)
DE      HLA-B*2702 CTL modulating peptide (B2702.84-75/75-84(T)).
KW      Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW      immunosuppressant; graft versus host disorder; transplantation; therapy;
KW      class I MHC; HLA-B*2702.
OS      Synthetic.
PN      WO9526979-A1.
PD      12-OCT-1995.
PF      05-APR-1995; U04349.
PR      05-APR-1994; US-222851.
PA      (STRD ) UNIV LELAND STANFORD JUNIOR.
PI      Clayberger C, Krensky AM, Parham P;
DR      WPI; 95-358582/46.
PT      Extension of acceptance period of transplants from MHC unmatched
PT      donor hosts - using Class I B*5-84 MHC antigen of the recipient
PT      host
PS      Example 15; Page 36; 80pp; English.
CC      R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
CC      Class I major histocompatibility complex (MHC) antigens. This sequence

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CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.062;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 YRLRLRLDER 10
 DB 1 YRLAIRLNER 10
 RESULT 6
 ID R92911 standard; peptide; 20 AA.
 AC R92911;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.84-75/84-75).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN WO9526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host

PS Example 15; Page 36; 80pp; English.
 PS R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is a dimer of residues 75-84 of the alpha-1 domain of the class I MHC
 CC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.062;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 YRLRLRLDER 10
 DB 1 YRLAIRLNER 10
 RESULT 7
 ID R92907 standard; peptide; 20 AA.
 AC R92907;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.84-75/84-75).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;

KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN WO9526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host

PS Example 15; Page 36; 80pp; English.
 PS R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.062;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 YRLRLRLDER 10
 DB 1 YRLAIRLNER 10
 RESULT 8
 ID R95428 standard; peptide; 20 AA.
 AC R95428;
 DT 12-NOV-1996 (first entry)
 DE HLA-B2702 84-75-84 palindromic.
 KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
 KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
 KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
 KW cytolysis; antigen presenting cell.
 OS Synthetic.
 PN WO9513288-A1.
 PD 18-MAY-1995.
 PF 10-NOV-1993; US-150493.
 PR 10-NOV-1994; U12985.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI: 95-194027/25.

PT Compsns. comprising lymphoid surface membrane proteins - which may
 PT inhibit cytolytic activity and differentiation of CTLs.
 PS Example; Page 12; 29pp; English.
 CC R95413, and R95415-R95431 represent palindromes and fragments of
 CC human-leucocyte-associated antigens. This sequence represents the
 CC HLA-B2702 84-75-84 palindromic. These sequences can be used to isolate
 CC the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
 CC protein associated with T-cell activation in mammalian T-cells, and is
 CC also immunologically cross reactive with the heat shock protein Hsc70.
 CC p74 is found in a limited number of cell types, but is particularly
 CC expressed on B and T cells. p74 can be isolated by lysis of a suitable
 CC cell with an amphoteric detergent, and then passed through an affinity
 CC column containing a covalently bound HLA-B2702 palindromic peptide.
 CC Compositions comprising the extracellular fragment of p74 combined with
 CC HLA-B2702.60-84 (see R95416), induces calcium influx, and inhibits
 CC cytotoxic T lymphocyte (CTL) differentiation or cytotoxicity. Candidate
 CC compounds can be screened for their effect on the cytolytic activity of
 CC T-cells, by combining them with the extracellular portion of p74 and
 CC determining the amount of binding between the candidate compound and p74.

CC Modulation of CTL activity can be inhibited in a cellular composition
 CC containing T-cells and antigen presenting cells (APCs), by adding to the
 CC mix the extracellular portion of p74, in an amount sufficient to compete
 CC with p74 for the binding of the p74 ligand.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.062;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDR 10
 ||| |||:||
 Db 1 YRLAIRLNER 10

RESULT 9

W33778
 ID W33778 standard; peptide: 20 AA.
 AC W33778;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #1.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplacantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 16; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is
 CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alfa1 domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.062;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDR 10
 ||| |||:||
 Db 1 YRLAIRLNER 10

RESULT 10

W33779
 ID W33779 standard; peptide: 20 AA.
 AC W33779;

DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #2.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplacantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 16; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is
 CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alfa1 domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
 Best Local Similarity 80.0%; Pred. No. 0.062;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLIRLDR 10
 ||| |||:||
 Db 1 YRLAIRLNER 10

RESULT 11

W33792
 ID W33792 standard; peptide: 20 AA.
 AC W33792;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2708.84-75/75-84T tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplacantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has

immunomodulating activity, including the N-terminal acylated and/or C-terminal amidated or esterified forms of up to 60 amino acids, where the peptide-type compound comprises the formula: A-B, where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino acid. The sequence in the brackets may optionally be absent or truncated at any peptide type bond within the brackets. The compounds comprise amino acid sequences related to a Class I HLA-B alpha domain (positions 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTLs) from undesirably attacking cells in a host or in vitro. They can also be used in combination with antigenic peptides or proteins of interest to activate CTLs. They can also inhibit the proliferation of T cells in response to anti-CD3. The peptide can be used for preventing rejection of transplants or for treating autoimmune diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus. The products can also be used for detection and diagnosis.

SQ Sequence 20 AA;

Query Match 79.6%; Score 39; DB 1; Length 20;
Best Local Similarity 80.0%; Pred. No. 0.062;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLRLDER 10
||| |||:||
Db 1 YRLATRLNER 10

RESULT 12

R92910 ID R92910 standard; peptide; 20 AA.
AC R92910;
DE 16-MAY-1996 (first entry)
DE HLA-B*2702 CTL modulating peptide (B2702.84-75(T)/75-84(T)).
KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW immunosuppressant; graft versus host disorder; transplantation; therapy;
KW class I MHC; HLA-B*2702.
OS Synthetic.
PN W09526979-A1.
PD 12-OCT-1995.
PF 05-APR-1995; U04349.
PR 05-APR-1994; US-222851.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM, Parham P;
DR WPI; 95-358582/46.
PT Extension of acceptance period of transplants from MHC unmatched donor hosts - using Class I B*5-84 MHC antigen of the recipient host
PT host

PS Example 15; Page 36; 80pp; English.
CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of Class I major histocompatibility complex (MHC) antigens. This sequence is an inverted dimer of residues 75-84 of the alpha-1 domain of the class I MHC HLA-B*2702. These sequences can be used to extend the period of acceptance by a recipient of a transplant from an MHC unmatched donor. The peptides are administered to a patient in conjunction with a subtherapeutic amount of an immunosuppressant. This is administered to the patient for a limited period of time (compared to the lifetime administration for current treatments). The peptides particularly modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs) of the patient.
SQ Sequence 20 AA;

Query Match 59.4%; Score 34; DB 1; Length 20;
Best Local Similarity 70.0%; Pred. No. 0.67;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLRLRLDER 10
||| |||:||
Db 1 YRLATRLNER 10

RESULT 13

R92908 ID R92908 standard; peptide; 20 AA.
AC R92908;
DE 16-MAY-1996 (first entry)
DE HLA-B*2702 CTL modulating peptide (B2702.84-75(T)/75-84).
KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW immunosuppressant; graft versus host disorder; transplantation; therapy;
KW class I MHC; HLA-B*2702.
OS Synthetic.
PN W09526979-A1.
PD 12-OCT-1995.
PF 05-APR-1995; U04349.
PR 05-APR-1994; US-222851.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM, Parham P;
DR WPI; 95-358582/46.
PT Extension of acceptance period of transplants from MHC unmatched donor hosts - using Class I B*5-84 MHC antigen of the recipient host
PT host

PS Example 15; Page 36; 80pp; English.
CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of Class I major histocompatibility complex (MHC) antigens. This sequence is an inverted dimer of residues 75-84 of the alpha-1 domain of the class I MHC HLA-B*2702. These sequences can be used to extend the period of acceptance by a recipient of a transplant from an MHC unmatched donor. The peptides are administered to a patient in conjunction with a subtherapeutic amount of an immunosuppressant. This is administered to the patient for a limited period of time (compared to the lifetime administration for current treatments). The peptides particularly modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs) of the patient.
SQ Sequence 20 AA;

Query Match 59.4%; Score 34; DB 1; Length 20;
Best Local Similarity 70.0%; Pred. No. 0.67;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLRLRLDER 10
||| |||:||
Db 1 YRLATRLNER 10

RESULT 14

R95430 ID R95430 standard; peptide; 20 AA.
AC R95430;
DE 12-NOV-1996 (first entry)
DE HLA-B*2702 84-75p/75-84p palindromic.
KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN W09513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI; 95-194027/25.
PT Compns. comprising lymphoid surface membrane proteins - which may inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 12; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of human-leucocyte-associated antigens. This sequence represents the HLA-B*2702 84-75p/75-84p palindromic. These sequences can be used to isolate the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane protein associated with T-cell activation in mammalian T-cells, and is also immunologically cross reactive with the heat shock protein Hsc70. p74 is found in a limited number of cell types, but is particularly expressed on B and T cells. p74 can be isolated by lysis of

CC a suitable cell with an amphoteric detergent, and then passed through an
 CC affinity column containing a covalently bound HLA-B2702 palindromic
 CC peptide. Compositions comprising the extracellular fragment of p74
 CC combined with HLA-B2702.60-84 (see R95416), induces calcium influx, and
 CC inhibits cytotoxic T lymphocyte (CTL) differentiation or cytotoxicity.
 CC Candidate compounds can be screened for their effect on the cytolytic
 CC activity of T-cells, by combining them with the extracellular portion of
 CC p74 and determining the amount of binding between the candidate compound
 CC and p74. Modulation of CTL activity can be inhibited in a cellular
 CC composition containing T-cells and antigen presenting cells (APCs), by
 CC adding to the mix the extracellular portion of p74, in an amount
 CC sufficient to compete with p74 for the binding of the p74 ligand.
 SQ Sequence 20 AA;

Query Match 69.4%; Score 34; DB 1; Length 20;
 Best Local Similarity 77.8%; Pred. No. 0.67;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLRLDE 9
 DB 1 YRLRLDE 9

RESULT 15

W33791
 ID W33791 standard; peptide; 20 AA.
 AC W33791;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702.84-757/75-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI; 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 4ipp: English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha1 domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 20 AA;

Query Match 69.4%; Score 34; DB 1; Length 20;
 Best Local Similarity 70.0%; Pred. No. 0.67;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLRLDER 10

DB 1 YRLATRLNER 10

Search completed: February 8, 2000, 01:29:38
 Job time: 1750 sec

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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:24 ; Search time 117.7 Seconds
(without alignments)
4.008 Million cell updates/sec

Title: US-08-653-294-13
Perfect score: 49
Sequence: 1 YRLAIRLDE 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : PIR_62.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	34	69.4	1154	2 S43275	hypothetical prote
2	34	69.4	1154	2 S43277	hypothetical prote
3	33	67.3	321	2 C64941	hypothetical prote
4	33	67.3	554	1 F70548	probable mend prot
5	32	65.3	151	2 C71113	probable frxa prot
6	32	65.3	185	2 S74416	hypothetical prote
7	32	65.3	318	2 A55429	11-cis retinol deh
8	32	65.3	319	2 I45845	11-cis retinol deh
9	32	65.3	349	1 RGEGL	nitrogen regulatio
10	32	65.3	349	2 A24114	probable acetate--
11	32	65.3	464	2 D42902	kynureninase (EC 3
12	32	65.3	464	2 S59898	heat-shock protein
13	32	65.3	546	2 S61294	DNA gyrase chain A
14	32	65.3	905	2 H71731	sugar-phosphate al
15	31	63.3	236	2 B72299	hypothetical prote
16	31	63.3	333	2 T05643	probable seryl-trn
17	31	63.3	463	2 B72500	glycoprotein gp13
18	31	63.3	485	1 B45343	probable UDP-N-ace
19	31	63.3	510	2 A70580	carboxylesterase (
20	31	63.3	544	2 B34089	DNA-directed RNA p
21	31	63.3	654	2 S58820	probable membrane
22	31	63.3	705	2 S54521	ethylene receptor
23	31	63.3	741	2 T16992	cation efflux syst
24	31	63.3	1063	2 A33830	cadmium, zinc, cob
25	31	63.3	1063	2 JC4700	zinc-finger protei
26	31	63.3	1214	2 JC2069	surface layer prot
27	31	63.3	1524	2 S68553	complement compone
28	31	63.3	1699	2 T14074	probable spindle p
29	31	63.3	2067	2 A42854	conserved hypothet
30	30	61.2	124	2 E70008	

31	30	61.2	125	2 D72544	hypothetical prote
32	30	61.2	141	1 F70457	hypothetical prote
33	30	61.2	204	2 C40899	hypothetical prote
34	30	61.2	252	2 H64752	probable transcrip
35	30	61.2	303	2 H71277	probable DNA adeni
36	30	61.2	312	2 T15371	hypothetical prote
37	30	61.2	336	2 H70693	hypothetical prote
38	30	61.2	465	2 JE0369	histidine acid pho
39	30	61.2	506	2 S37583	RING finger protei
40	30	61.2	513	1 TVHURF	ret finger protein
41	30	61.2	540	2 S76869	hypothetical prote
42	30	61.2	545	2 T00485	probable phosphori
43	30	61.2	547	2 A56575	puff-specific nucl
44	30	61.2	610	2 G69130	conserved hypothet
45	30	61.2	723	2 T14605	probable cell divi

ALIGNMENTS

RESULT 1

S43275

hypothetical protein 2 - Neurospora crassa retrotransposon Tad1-1

C:Species: Neurospora crassa

C>Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 09-Sep-1997

C:Accession: S43275

R:Cambareri, E.B.; Helber, J.; Kinsey, J.A.

Mol. Gen. Genet. 242, 658-665, 1994

A:Title: Tad1-1, an active LINE-like element of Neurospora crassa.

A:Reference number: S43274; MUID:94203179

A:Accession: S43275

A:Molecule type: DNA

A:Residues: 1-1154 <CAM>

A:Cross-references: EMBL:L255662; NID:g409759; PID:g409761

C:Genetics:

A:Mobile element: retrotransposon Tad1-1

Query Match 69.4%; Score 34; DB 2; Length 1154;
Best Local Similarity 66.7%; Pred. No. 51;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9

DB 1136 YRLAVELEE 1144

RESULT 2

S43277

hypothetical protein 2 - Neurospora crassa retrotransposon Tad3-2

C:Species: Neurospora crassa

C>Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 09-Sep-1997

C:Accession: S43277

R:Cambareri, E.B.; Helber, J.; Kinsey, J.A.

Mol. Gen. Genet. 242, 658-665, 1994

A:Title: Tad1-1, an active LINE-like element of Neurospora crassa.

A:Reference number: S43274; MUID:94203179

A:Accession: S43277

A:Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-1154 <CAM>

A:Cross-references: EMBL:L255663; NID:g409762; PID:g409764

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, November 1993

Query Match 69.4%; Score 34; DB 2; Length 1154;
Best Local Similarity 66.7%; Pred. No. 51;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9

DB 1136 YRLAVELEE 1144

```

RESULT 3
C64941
hypothetical protein b1803 - Escherichia coli (strain K-12)
C:Species: Escherichia coli
C>Date: 12-Sep-1997 #sequence_revision 17-Sep-1997 #text_change 29-Sep-1999
C:Accession: C64941
R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; Cohen, A.; Rose, D.J.; Mau, B.; Shao, Y.
Science 277, 1453-1462, 1997
A:Title: The complete genome sequence of Escherichia coli K-12.
A:Reference number: A64720; MUID:97426617
A:Accession: C64941
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-321 <BLAT>
A:Cross-references: GB:AE000274; GB:U00096; NID:g1788089; PIDN:AACT4873.1; PID:g1788104;
A:Experimental source: strain K-12, substrain MG1655
C:Superfamily: phthalate dioxygenase reductase; cytochrome-b5 reductase homology; ferredoxin
F:11-225/Domain: cytochrome-b5 reductase homology <CBR>
F:254-309/Domain: ferredoxin [2Fe-2S] homology <FER>

Query Match 67.3%; Score 33; DB 2; Length 321;
Best Local Similarity 55.6%; Pred. No. 22;
Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Qy 1 YRLAIRLDE 9
|||:|:|:|
Db 70 YQIAVRLEE 78

RESULT 4
F70548
probable menD protein - Mycobacterium tuberculosis (strain H37RV)
C:Species: Mycobacterium tuberculosis
C>Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 10-Sep-1999
C:Accession: F70548
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon, S.; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holtroyd, S.; Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.;
Nature 393, 537-544, 1998
A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete genome
A:Reference number: A70500; MUID:98295987
A:Accession: F70548
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-554 <COL>
A:Cross-references: GB:295558; GB:AL123456; NID:g3261781; PID:e316800; PID:g2114017
A:Experimental source: strain H37RV
C:Genetics:
A:Gene: menD
C:Superfamily: menD protein

Query Match 67.3%; Score 33; DB 1; Length 554;
Best Local Similarity 66.7%; Pred. No. 39;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLAIKRLD 10
||:|:|:|
Db 48 RLHVRIDER 56

RESULT 5
C71113
probable frxA protein - Pyrococcus horikoshii
C:Species: Pyrococcus horikoshii
C>Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 14-Aug-1998
C:Accession: C71113
R:Kawarabayashi, Y.; Sawada, M.; Horikawa, H.; Haikawa, Y.; Hino, Y.; Yamamoto, S.; Sekiguchi, M.; Ohfuku, Y.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; Kushida, N.; Oguchi

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DNA Res. 5, 55-76, 1998
A:Title: Complete sequence and gene organization of the genome of a hyper-thermophilic
A:Reference number: A71000; MUID:98344137
A:Accession: C71113
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-151 <KAW>
A:Cross-references: GB:AF000003; NID:g3236130; PID:d1030708; PID:g3257082
A:Experimental source: strain OT3
A:Note: this accession replaces an interim accession for a sequence replaced by GenBank
C:Genetics:
A:Gene: PH0674

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Query Match 65.3%; Score 32; DB 2; Length 151;
Best Local Similarity 77.8%; Pred. No. 16;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 RLAIKRLD 10
||:|:|:|
Db 96 RLIIELDER 104

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RESULT 6
S74416
hypothetical protein s110687 - Synecocystis sp. (strain PCC 6803)
C:Species: Synecocystis sp.
A:Variety: PCC 6803
C>Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 21-Aug-1998
C:Accession: S74416
R:Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, E.; Nakamura, Y.; Miyajima, O. K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yasuda, M.
DNA Res. 3, 109-136, 1996
A:Title: Sequence analysis of the genome of the unicellular cyanobacterium Synecocystis
S:Reference number: S74322; MUID:97061201
A:Accession: S74416
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-185 <KAN>
A:Cross-references: EMBL:D64001; GB:AB001339; NID:g1001102; PID:d1010985; PID:g1001119
A:Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996

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Query Match 65.3%; Score 32; DB 2; Length 185;
Best Local Similarity 60.0%; Pred. No. 20;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 1 YRLAIRLDER 10
|||:|:|:|
Db 35 YRLAIRILQR 44

```

```

RESULT 7
A55429
11-cis retinol dehydrogenase (EC 1.1.1.-) - bovine
C:Species: Bos primigenius taurus (cattle)
C>Date: 10-Feb-1995 #sequence_revision 10-Feb-1995 #text_change 29-Sep-1999
C:Accession: A55429
R:Simon, A.; Hellman, U.; Wernstedt, C.; Eriksson, U.
J. Biol. Chem. 270, 1107-1112, 1995
A:Title: The retinal pigment epithelial-specific 11-cis retinol dehydrogenase belongs
A:Reference number: A55429; MUID:95138097
A:Accession: A55429
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: mRNA
A:Residues: 1-318 <SIM>
A:Cross-references: GB:X82262; NID:g663170; PIDN:CAA57715.1; PID:g663171
C:Superfamily: retinol dehydrogenase; short-chain alcohol dehydrogenase homology
C:Keywords: membrane protein; NAD; oxidoreductase
F:29-206/Domain: short-chain alcohol dehydrogenase homology <SADH>

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Query Match 65.3%; Score 32; DB 2; Length 318;
 Best Local Similarity 75.0%; Pred. No. 35;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDER 10
 ||:||||:|
 Db 44 LAIRLDQR 51

RESULT 8

I45845
 11-cis-retinol dehydrogenase (EC 1.1.1.1-) - bovine
 C:Species: Bos primigenius taurus (cattle)
 C:Date: 15-Oct-1996 #sequence_revision 15-Oct-1996 #text_change 29-Sep-1999
 C:Accession: I45845
 R:Driessen, C.A.; Janssen, B.P.; Winkens, H.J.; van Vugt, A.H.; de Leeuw, T.L.; Janssen, Invest. Ophthalmol. Vis. Sci. 36, 1988-1996, 1993
 A:Title: Cloning and expression of a cDNA encoding bovine retinal pigment epithelial 11-
 A:Reference number: I45845; MUID:95386398
 A:Accession: I45845
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-319 <DRI>
 A:Cross-references: GB:I36533; NID:g1054530; PIDN:AAA80694.1; PID:g1054531
 C:Superfamily: retinol dehydrogenase; short-chain alcohol dehydrogenase homology
 C:Keywords: oxidoreductase
 F:30-207/Domain: short-chain alcohol dehydrogenase homology <SADH>

Query Match 65.3%; Score 32; DB 2; Length 319;
 Best Local Similarity 75.0%; Pred. No. 35;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDER 10
 ||:||||:|
 Db 45 LAIRLDQR 52

RESULT 9

RGEGL
 nitrogen regulation protein II (EC 2.7.3.-) ntrB - Escherichia coli
 N:Alternate names: regulatory protein glnL
 C:Species: Escherichia coli
 C:Date: 31-Mar-1990 #sequence_revision 31-Mar-1990 #text_change 16-Jul-1999
 C:Accession: A30377; S40814; B23970; H65191; A39765; Q00553
 R:Miranda-Rios, J.; Sanchez-Pescador, R.; Urdea, M.; Covarrubias, A.A.
 Nucleic Acids Res. 15, 2757-2770, 1987
 A:Title: The complete nucleotide sequence of the glnALG operon of Escherichia coli K12.
 A:Reference number: A30377; MUID:87174797
 A:Accession: A30377
 A:Molecule type: DNA
 A:Residues: 1-349 <MR>
 A:Cross-references: EMBL:X05173; NID:g41562; PIDN:CAA28807.1; PID:g41564
 A:Experimental source: K-12
 R:Plunkett III, G.; Burland, V.; Daniels, D.L.; Blattner, F.R.
 Nucleic Acids Res. 21, 3391-3398, 1993
 A:Title: Analysis of the Escherichia coli genome. III. DNA sequence of the region from 8
 A:Reference number: S40802; MUID:93347969
 A:Accession: S40814
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-349 <PLU>
 A:Cross-references: EMBL:I19201; NID:g304961; PIDN:AA03003.1; PID:g304974
 A:Experimental source: strain K-12, substrain MG1655
 A:Note: the nucleotide sequence was submitted to the EMBL Data Library, October 1993
 R:Roche, M.; Vazquez, M.; Garcia-Rubio, A.; Covarrubias, A.A.
 Gene 37, 91-99, 1985
 A:Title: Nucleotide sequence of the glnA-glnL intercistronic region of Escherichia coli.
 A:Reference number: A91533; MUID:86031370
 A:Accession: B23970
 A:Molecule type: DNA
 A:Residues: 1-24 <ROC>
 A:Cross-references: GB:K02176; GB:M11581; NID:g146160; PIDN:AAA23881.1; PID:g146162

R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.;
 A.; Rose, D.J.; Mau, B.; Shao, Y.
 Science 277, 1453-1462, 1997
 A:Title: The complete genome sequence of Escherichia coli K-12.
 A:Reference number: A64720; MUID:97428617
 A:Accession: H65191
 A:Status: nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-349 <BLAT>
 A:Cross-references: GB:AE000462; GB:U00096; NID:g1790295; PIDN:AAC76866.1; PID:g17903
 A:Experimental source: strain K-12, substrain MG1655
 R:Ninfa, A.J.; Bennett, R.L.
 J. Biol. Chem. 266, 6888-6893, 1991
 A:Title: Identification of the site of autophosphorylation of the bacterial protein K
 A:Reference number: A39765; MUID:91201336
 A:Accession: A39765
 A:Status: preliminary
 A:Molecule type: protein
 A:Residues: 2-11;136-142;158-162,'X',164-169 <NIN>
 C:Genetics:
 A:Gene: glnL; ntrB
 A:Map position: 87 min
 C:Function:

A:Description: de-uridylylated P-II forms a complex with nitrogen regulation protein
 the uridylylated form of P-II does not complex with ntrB; free ntrB phosphorylates n
 A:Note: phosphorylated nitrogen regulation protein I (ntrC) activates transcription o
 C:Superfamily: glnL regulatory protein II; sensor histidine kinase homology
 C:Keywords: ATP; autophosphorylation; phosphohistidine; phosphoprotein; phosphotransf
 F:104-346/Domain: sensor histidine kinase homology <SHK>
 F:139/Binding site: phosphate (His) (covalent) (by autophosphorylation) #status predi
 F:329/Binding site: ATP (Lys) #status predicted

Query Match 65.3%; Score 32; DB 1; Length 349;
 Best Local Similarity 75.0%; Pred. No. 39;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLD 8
 |||||:|
 Db 277 YRLAARID 284

RESULT 10

A24114
 nitrogen regulation protein II (EC 2.7.3.-) ntrB - Klebsiella pneumoniae
 C:Species: Klebsiella pneumoniae
 C:Date: 22-Jul-1987 #sequence_revision 22-Jul-1987 #text_change 20-Aug-1999
 C:Accession: A24114
 R:MacFarlane, S.A.; Merrick, M.
 Nucleic Acids Res. 13, 7591-7607, 1985
 A:Reference number: A24114; MUID:86067184
 A:Accession: A24114
 A:Molecule type: DNA
 A:Residues: 1-349 <MAC>
 A:Cross-references: GB:X03146; NID:g43893; PIDN:CAA26923.1; PID:g43895
 C:Genetics:
 A:Gene: ntrB
 C:Superfamily: glnL regulatory protein II; sensor histidine kinase homology
 C:Keywords: ATP; autophosphorylation; phosphohistidine; phosphoprotein; phosphotransf
 F:104-346/Domain: sensor histidine kinase homology <SHK>
 F:139/Binding site: phosphate (His) (covalent) (by autophosphorylation) #status predi
 F:329/Binding site: ATP (Lys) #status predicted

Query Match 65.3%; Score 32; DB 2; Length 349;
 Best Local Similarity 75.0%; Pred. No. 39;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLD 8
 |||||:|
 Db 277 YRLAARID 284

RESULT 11

D42902
probable acetate--CoA ligase (EC 6.2.1.1) - Pseudomonas aeruginosa (fragment)
N:Alternate names: probable acetyl-CoA synthetase
C:Species: Pseudomonas aeruginosa
C:Date: 04-Mar-1993 #sequence_revision 24-Oct-1997 #text_change 24-Oct-1997
C:Accession: S27604; D42902
R:Steele, M.I.; Lorenz, D.; Hatter, K.; Parks, A.; Sokatch, J.R.
submitted to the EMBL Data Library, July 1992
A:Description: Characterization of the mmsAB operon of Pseudomonas aeruginosa PAO encoding met
A:Reference number: S27601
A:Accession: S27604
A:Molecule type: DNA
A:Residues: 1-464 <STE>
A:Cross-references: EMBL:M84911; NID:gl51360; PID:g551933
R:Steele, M.I.; Lorenz, D.; Hatter, K.; Park, A.; Sokatch, J.R.
J. Biol. Chem. 267, 13585-13592, 1992
A:Title: Characterization of the mmsAB operon of Pseudomonas aeruginosa PAO encoding met
A:Reference number: A42902; MUID:92317087
A:Accession: D42902
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-79 <ST>
A:Experimental source: PAO, ATCC 15692
A:Note: sequence extracted from NCBI backbone (NCBIN:107704, NCBI:P:107709)
C:Superfamily: acetate--CoA ligase homology
C:Keywords: acid-thiol ligase
F:105-464/Domain: acetate--CoA ligase homology (fragment) <ACL>

Query Match 65.3%; Score 32; DB 2; Length 464;
Best Local Similarity 55.0%; Pred. No. 52;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
QY 1 YRLAIRLDE 9
| | | | |
Db 146 YELALRID 154

RESULT 12

S59898
kynureninase (EC 3.7.1.3) - rat
C:Species: Rattus norvegicus (Norway rat)
C:Date: 20-Jul-1996 #sequence_revision 13-Mar-1997 #text_change 13-Mar-1997
C:Accession: S59898
R:Takeuchi, F.; Tsubouchi, R.; Yoshino, M.; Shibata, Y.
Biochim. Biophys. Acta 1252, 185-188, 1995
A:Title: Amino-acid sequence of rat liver kynureninase.
A:Reference number: S59898; MUID:96049498
A:Accession: S59898
A>Status: preliminary
A:Molecule type: protein
A:Residues: 1-464 <TAK>
C:Keywords: hydrolase

Query Match 65.3%; Score 32; DB 2; Length 464;
Best Local Similarity 75.0%; Pred. No. 52;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 2 RLAIRLDE 9
| | | | |
Db 28 RVALRLDE 35

RESULT 13

S61294
heat-shock protein - Pyrococcus sp.
C:Species: Pyrococcus sp.
C:Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 20-Aug-1999
C:Accession: S61294
R:Izawa, Y.; Kakiwara, H.; Takagi, M.; Imanaka, T.
submitted to the EMBL Data Library, March 1994

A:Description: Cloning and analysis of the heat shock protein gene from a new hyperthermophile
A:Reference number: S61294
A:Accession: S61294
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-546 <IZA>
A:Cross-references: EMBL:D29672; NID:g473964; PIDN:BAA06143.1; PID:d1006705; PID:g473
C:Superfamily: molecular chaperone t-complex-type
C:Keywords: heat shock

Query Match 65.3%; Score 32; DB 2; Length 546;
Best Local Similarity 100.0%; Pred. No. 61;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 3 LAIRLDE 9
| | | | |
Db 418 LAIRLDE 424

RESULT 14

H71731
DNA gyrase chain A (gyrA) RP206 - Rickettsia prowazekii
C:Species: Rickettsia prowazekii
C:Date: 21-Nov-1998 #sequence_revision 21-Nov-1998 #text_change 16-Jul-1999
C:Accession: H71731
R:Andersson, S.G.E.; Zomorodipour, A.; Andersson, J.O.; Sicheritz-Ponten, T.; Alsmark
Nature 396, 133-140, 1998
A:Title: The genome sequence of Rickettsia prowazekii and the origin of mitochondria.
A:Reference number: A71630; MUID:99039499
A:Accession: H71731
A>Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-905 <AND>
A:Cross-references: GB:AJ235270; GB:AJ235269; NID:g3860572; PIDN:CAA14671.1; PID:el34
A:Experimental source: strain Madrid E
C:Genetics:

Query Match 65.3%; Score 32; DB 2; Length 905;
Best Local Similarity 75.0%; Pred. No. 1e+02;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 2 RLAIRLDE 9
| | | | |
Db 665 RLAIRLDE 672

RESULT 15

B72299
sugar-phosphate aldolase - Thermotoga maritima (strain MSB8)
C:Species: Thermotoga maritima
C:Date: 11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 11-Jun-1999
C:Accession: B72299
R:Nelson, K.E.; Clayton, R.A.; Gill, S.R.; Gwinn, M.L.; Dodson, R.J.; Haft, D.H.; Hic
Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.A.; Richardson,
C.M.
Nature 399, 323-329, 1999
A:Title: Evidence for lateral gene transfer between Archaea and Bacteria from genome
A:Reference number: A72200; MUID:99287316
A:Accession: B72299
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-236 <ARN>
A:Cross-references: GB:AE001767; GB:AE000512; NID:g4981611; PID:g4981616; TIGR:TM1072
A:Experimental source: strain MSB8
C:Genetics:
A:Gene: TM1072

Query Match 63.3%; Score 31; DB 2; Length 236;
Best Local Similarity 62.5%; Pred. No. 42;
Matches 5; Conservative 3; Mismatches 0; Indels 0; Gaps 0;
Qy 3 LAIRLDER 10
:::|||||
Db 28 ISVRLDER 35

Search completed: February 7, 2000, 11:54:25
Job time: 24335 sec

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OM protein - protein search, using sw model

Run on: February 8, 2000, 00:59:52 ; Search time 63.71 seconds
(without alignments)
4.688 Million cell updates/sec

Title: US-08-653-294-13
Perfect score: 49
Sequence: 1 YRLAIRLDR 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 82229 seqs, 29864866 residues

Total number of hits satisfying chosen parameters: 82229

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : SwissProt_38:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	33	67.3	321	1 YEAX_ECOLI	P76254 escherichia
2	32	65.3	318	1 RDHL_BOVIN	Q27979 bos taurus
3	32	65.3	349	1 NTRB_ECOLI	P06712 escherichia
4	32	65.3	349	1 NTRB_KLEPN	P06718 klebsiella
5	32	65.3	464	1 KYNU_RAT	P70712 rattus norv
6	32	65.3	464	1 YWMS_PSEAE	P28812 pseudomonas
7	32	65.3	546	1 THS_PYRKO	Q52500 pyrococcus
8	32	65.3	548	1 THSA_THRKL	O24729 thermococcus
9	32	65.3	905	1 GIRA_RICPR	P41080 rickettsia
10	31	63.3	485	1 VGLC_HSVF4	P22596 equine herp
11	31	63.3	510	1 MURF_MYCTU	O06220 mycobacteri
12	31	63.3	544	1 ESTP_DROME	P18167 drosophila
13	31	63.3	654	1 RPC3_YEAST	P32349 saccharomyc
14	31	63.3	705	1 YM37_YEAST	Q03824 saccharomyc
15	31	63.3	1014	1 MMLB_MYCLE	O06079 mycobacteri
16	31	63.3	1063	1 CZCA_ALGCEU	P13511 alcaligenes
17	31	63.3	1063	1 CZCA_ALGSP	P94177 alcaligenes
18	31	63.3	1214	1 BR14_HUMAN	P52201 homo sapien
19	31	63.3	2067	1 BMBB_EMENI	P33144 emericella
20	30	61.2	252	1 YAGI_ECOLI	P77300 escherichia
21	30	61.2	303	1 DMA_TREPA	O33844 treponema p
22	30	61.2	513	1 RFP_HUMAN	P14373 homo sapien
23	30	61.2	522	1 RFP_MOUSE	Q62158 mus musculu
24	30	61.2	547	1 BX42_DROME	P39736 drosophila
25	30	61.2	663	1 TERM_ADEB3	O55439 bovine aden
26	30	61.2	1176	1 NLR_NEUCR	P38681 neurospora
27	30	61.2	3491	1 ERYL_SACER	Q03131 saccharopol
28	30	61.2	3587	1 SRFL_BACSU	P27206 bacillus su
29	29	59.2	108	1 YCT9_YEAST	P39534 saccharomyc
30	29	59.2	158	1 YCBM_BACSU	P42245 bacillus su
31	29	59.2	189	1 VH02_VACCC	P20496 vaccinia vi
32	29	59.2	189	1 VH02_VACCV	P08583 vaccinia vi
33	29	59.2	189	1 VH02_VARV	P33061 variola vir
34	29	59.2	210	1 FLPA_ARCFU	O28192 archaeoglob

RESULT 1

YEAX_ECOLI
ID YEAX_ECOLI STANDARD; PRT; 321 AA.
AC P76254; O07972; O07970;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE PUTATIVE DIOXYGENASE BETA SUBUNIT YEAX (EC 1.-.-.-).
GN YEAX.
OS Escherichia coli.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Escherichia.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=K12 / MG1655;
RX MEDLINE; 97426617.
RA BLATTNER F. R., PLUNKETT G. III, BLOCH C.A., PERNA N.T., BURLAND V.,
RILEY M., COLLADO-VIDES J., GLASNER F.D., RODE C.K., MAYHEW G.F.,
GREGOR J., DAVIS N.W., KIRKPATRICK H.A., GOEDEN M.A., ROSE D.J.,
MAU B., SHAO Y.;
RT "The complete genome sequence of Escherichia coli K-12.";
RL Science 277:1453-1474(1997).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=K12;
RX MEDLINE; 97251358.
RA ITOH T., AIBA H., BABA T., FUJITA K., HAYASHI K., INADA T., ISONO K.,
KASAI H., KIMURA S., KITAKAWA M., KITAGAWA M., MAKINO K., MIKI T.,
MIZOBUCHI K., MORI H., MORI T., MOTOMURA K., NAKADE S., NAKAMURA Y.,
NASHIMOTO H., NISHIO Y., OSHIMA T., SAITO N., SAMPEI G., SEKI Y.,
SIVASUNDARAM S., TAGAMI H., TAKEDA J., TAKEMOTO K., WADA C.,
YAMAMOTO Y., HORIUCHI T.;
RT "A 460-kb DNA sequence of the Escherichia coli K-12 genome
corresponding to the 40.1-50.0 min region on the linkage map.";
RL DNA Res. 3:379-392(1996).
CC -1- COFACTOR: FMN (BY SIMILARITY).
CC -1- SUBUNIT: PROBABLE HETERODIMER OF YEAW AND YEAX.
CC -1- SIMILARITY: IN THE C-TERMINAL, BELONGS TO THE 2FE2S PLANT-TYPE
FERREDOXIN FAMILY.
CC -1- SIMILARITY: BELONGS TO THE PDR/VANB FAMILY.
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CC
CC EMBL; AE000274; AAC74873.1; -.
CC DR EMBL; D90824; CAB21531.1; -.
CC DR EMBL; D90823; CAB21524.1; -.
CC DR HSSP; P33164; 2PIA.
CC DR ECGENE; EG13510; YEAX.
CC DR PROSITE; PS00197; 2FE2S_FERREDOXIN; 1.
CC DR PFAM; PF00111; fer2; 1.
CC DR PFAM; PF00175; oxidored_fad; 1.

ALIGNMENTS

35 29 59.2 246 1 TRYP_MOUSE
36 29 59.2 279 1 LEP3_ERWCA
37 29 59.2 294 1 RL5A_SCHPO
38 29 59.2 294 1 RL5B_SCHPO
39 29 59.2 297 1 RL5_HELAN
40 29 59.2 299 1 RL5_BOMMO
41 29 59.2 306 1 MK16_YEAST
42 29 59.2 334 1 RUVB_THEMA
43 29 59.2 377 1 CAHL_CHLRE
44 29 59.2 409 1 METK_SYNY3
45 29 59.2 428 1 YURL_YEAST
P07146 mus musculu
P31712 erwinia car
P52822 schizosacch
O74306 schizosacch
O65333 helianthus
O76190 bombyx mori
P10962 saccharomyc
Q56313 thermotoga
P20507 chlamydomon
P72871 synechocyst
P26725 saccharomyc

KW Hypothetical protein; Oxidoreductase; Flavoprotein; FMN; NAD;
 Iron-sulfur; Electron transport.
 FT NP_BIND 6 103 FMN (BY SIMILARITY).
 FT NP_BIND 113 226 NAD (BY SIMILARITY).
 FT METAL 270 270 IRON-SULFUR (2FE-2S) (BY SIMILARITY).
 FT METAL 275 275 IRON-SULFUR (2FE-2S) (BY SIMILARITY).
 FT METAL 278 278 IRON-SULFUR (2FE-2S) (BY SIMILARITY).
 FT METAL 309 309 IRON-SULFUR (2FE-2S) (BY SIMILARITY).
 SQ SEQUENCE 321 AA: 35661 MW; 9E85CC68 CRC32;

Query Match 67.3%; Score 33; DB 1; Length 321;
 Best Local Similarity 55.6%; Pred. No. 11;
 Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9
 |::|::|::|
 Db 70 YQIAVRLEE 78

RESULT 2
 RDH1_BOVIN
 ID RDH1_BOVIN STANDARD; PRT; 318 AA.
 AC Q27979;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DE 11-CIS RETINOL DEHYDROGENASE (EC 1.1.1.105) (11-CIS RDH) (P32).
 GN RDH1.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Cranialata; Vertebrata; Mammalia;
 OC Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OC Bovinae; Bos.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE-EYE;
 RX MEDLINE: 95138097.
 RA SIMON A., HELLMAN U., WERNSTEDT C., ERIKSSON U.;
 RT "The retinal pigment epithelial-specific 11-cis retinol dehydrogenase
 belongs to the family of short chain alcohol dehydrogenases.";
 RL J. Biol. Chem. 270:1107-1112(1995).
 CC -!- FUNCTION: STEREOSPECIFIC 11-CIS RETINOL DEHYDROGENASE, WHICH
 CATALYZES THE FINAL STEP IN THE BIOSYNTHESIS OF 11-CIS
 RETINALDEHYDE, THE UNIVERSAL CHROMOPHORE OF VISUAL PIGMENTS.
 CC ACTIVE IN THE PRESENCE OF NAD+ AS COFACTOR BUT NOT IN THE PRESENCE
 OF NADP.
 CC -!- CATALYTIC ACTIVITY: RETINOL + NAD(+) -> RETINAL + NADH.
 CC -!- PATHWAY: CATALYZES THE PRIMARY AND RATE-LIMITING STEP IN RETINOIC
 ACID SYNTHESIS.
 CC -!- SUBCELLULAR LOCATION: MEMBRANE-ASSOCIATED.
 CC -!- SIMILARITY: BELONGS TO THE SHORT-CHAIN DEHYDROGENASES/REDUCTASES
 FAMILY (SDR).
 CC
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 CC
 CC EMBL: X82262; CAA57715.1; -
 DR HSP: P14061; 1FDW.
 DR PROSITE: PS00061; ADH_SHORT; FALSE_NEG.
 DR PFAM: PF00106; adh_short; 1.
 KW Oxidoreductase; NAD. 56 NADP (BY SIMILARITY).
 FT NP_BIND 32 175 BY SIMILARITY.
 FT ACT_SITE 175 175
 SQ SEQUENCE 318 AA: 35036 MW; 7022A583 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 318;
 Best Local Similarity 75.0%; Pred. No. 18;

Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 QY 3 LAIRLDER 10
 |::|::|::|
 Db 44 LAIRLDQR 51
 RESULT 3
 NTRB_ECOLI
 ID NTRB_ECOLI STANDARD; PRT; 349 AA.
 AC P06712;
 DT 01-JAN-1988 (Rel. 06, Created)
 DT 01-JAN-1988 (Rel. 06, Last sequence update)
 DT 01-NOV-1997 (Rel. 35, Last annotation update)
 DE NITROGEN REGULATION PROTEIN NR(II) (EC 2.7.3.-).
 GN GLNL OR NTRB OR GLNR.
 OS Escherichia coli.
 OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 OC Escherichia.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-K12;
 RX MEDLINE: 87174797.
 RA MIRANDA-RIOS J., SANCHEZ-PESCADOR R., URDEA M., COVARRUBIAS A.A.;
 RT "The complete nucleotide sequence of the glnALG operon of Escherichia
 coli K12.";
 RL Nucleic Acids Res. 15:2757-2770(1987).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-K12 / MG1655;
 RX MEDLINE: 93347969.
 RA PLUNKETT G. III, BURLAND V.D., DANIELS D.L., BLATTNER F.R.;
 RT "Analysis of the Escherichia coli genome. III. DNA sequence of the
 region from 87.2 to 89.2 minutes.";
 RL Nucleic Acids Res. 21:3391-3398(1993).
 RN [3]
 RP SEQUENCE OF 1-22 FROM N.A.
 RX MEDLINE: 85006814.
 RA UENO-NISHIO S., MANGO S., REITZER L.J., MAGASANIK B.;
 RT "Identification and regulation of the glnL operator-promoter of the
 complex glnALG operon of Escherichia coli.";
 RL J. Bacteriol. 160:379-384(1984).
 RN [4]
 RP SEQUENCE OF 1-24 FROM N.A.
 RX MEDLINE: 86031370.
 RA ROCHA M., VAZQUEZ M., GARCIA-RUBIO A., COVARRUBIAS A.A.;
 RT "Nucleotide sequence of the glnA-glnL intercistronic region of
 Escherichia coli.";
 RL Gene 37:91-99(1985).
 RN [5]
 RP PHOSPHORYLATION SITE.
 RX MEDLINE: 91201336.
 RA NINFA A.J., BENNETT R.L.;
 RT "Identification of the site of autophosphorylation of the bacterial
 protein kinase/phosphatase NR11.";
 RL J. Biol. Chem. 266:6888-6893(1991).
 CC -!- FUNCTION: NTRB ACTS AS A SIGNAL TRANSDUCER WHICH RESPONDS TO THE
 NITROGEN LEVEL OF CELL AND MODULATES THE ACTIVITY OF NTRC. IN
 NITROGEN LIMITATION NTRB ACTIVATES NTRC BY PHOSPHORYLATING IT.
 CC WHILE IN NITROGEN EXCESS NTRC IS DEPHOSPHORYLATED AND CONSEQUENTLY
 INACTIVATED BY NTRB.
 CC -!- SIMILARITY: TO OTHER PROKARYOTIC SENSORY TRANSDUCTION HISTIDINE
 KINASES.
 CC
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 CC
 CC EMBL: X05173; CAA28807.1; -

DR EMBL; K02176; AAA23881.1; --
 DR EMBL; L19201; AAB03003.1; --
 DR EMBL; AE000462; AAC76866.1; --
 DR PIR; Q00553; RGECLG.
 DR PIR; B23970; B23970.
 DR PIR; S40814; S40814.
 DR ECGENE; EG10387; GLNL.
 DR PFAM; PF00512; signal; 1.
 KW Sensory transduction; Transferase; Kinase; Phosphorylation;
 KW Nitrogen fixation; ATP-binding.
 FT DOMAIN 116 349 TRANSMITTER DOMAIN (POTENTIAL).
 FT MOD_RES 139 139 PHOSPHORYLATION (AUTO-).
 FT BINDING 329 329 ATP (BY SIMILARITY).
 SQ SEQUENCE 349 AA; 38556 MW; 6A017919 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 349;
 Best Local Similarity 75.0%; Pred. No. 19;
 Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 YRLAIRLD 8
 |||||
 Db 277 YRLAARID 284

RESULT 4
 NTRB_KLEPN ID NTRB_KLEPN STANDARD; PRT; 349 AA.
 AC P06218;
 DT 01-JAN-1988 (Rel. 06, Created)
 DT 01-JAN-1988 (Rel. 06, Last sequence update)
 DT 01-OCT-1994 (Rel. 30, Last annotation update)
 DE NITROGEN REGULATION PROTEIN NTRB (EC 2.7.3.-).
 GN NTRB.
 OS Klebsiella pneumoniae.
 OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 OC Klebsiella.
 RN SEQUENCE FROM N.A.
 RP MEDLINE; 86067184.
 RA MCFARLANE S.A., MERRICK M.J.;
 RT "The nucleotide sequence of the nitrogen regulation gene ntrB and the
 glnA-ntrB intergenic region of Klebsiella pneumoniae.";
 RL Nucleic Acids Res. 13:7591-7606(1985).
 CC -1- FUNCTION: NTRB ACTS AS A SIGNAL TRANSDUCER WHICH RESPONDS TO THE
 NITROGEN LEVEL OF CELL AND MODULATES NTRC BY PHOSPHORYLATING IT.
 CC NITROGEN LIMITATION NTRB ACTIVATES NTRC BY PHOSPHORYLATING IT.
 CC WHILE IN NITROGEN EXCESS NTRC IS DEPHOSPHORYLATED AND CONSEQUENTLY
 CC INACTIVATED BY NTRB.
 CC -1- SIMILARITY: TO OTHER PROKARYOTIC SENSORY TRANSDUCTION HISTIDINE
 KINASES.
 CC
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 CC
 DR EMBL; X03146; CAA26923.1; --
 DR PIR; A24114; A24114.
 DR PFAM; PF00512; signal; 1.
 KW Sensory transduction; Transferase; Kinase; Phosphorylation;
 KW Nitrogen fixation; ATP-binding.
 FT DOMAIN 116 349 TRANSMITTER DOMAIN (POTENTIAL).
 FT MOD_RES 139 139 PHOSPHORYLATION (AUTO-) (BY SIMILARITY).
 FT BINDING 329 329 ATP (BY SIMILARITY).
 SQ SEQUENCE 349 AA; 38409 MW; 4BAC1813 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 349;
 Best Local Similarity 75.0%; Pred. No. 19;

Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 YRLAIRLD 8
 |||||
 Db 277 YRLAARID 284

RESULT 5
 KYNU_RAT ID KYNU_RAT STANDARD; PRT; 464 AA.
 AC P70712;
 DT 15-DEC-1998 (Rel. 37, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 15-DEC-1998 (Rel. 37, Last annotation update)
 DE KYNURENINASE (EC 3.7.1.3) (L-KYNURENINE HYDROLASE).
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 RN [1]
 RP SEQUENCE FROM N.A., AND PARTIAL SEQUENCE.
 RC TISSUE=LIVER;
 RX MEDLINE; 97324088.
 RA TOMA S., NAKAMURA M., TONE S., OKUNO E., KIDO R., BRETON J.,
 RA AVANZI N., COZZI L., SPECIALE C., MOSTARDINI M., GATTI S., BENATTI L.;
 RT "Cloning and recombinant expression of rat and human kynureninase.";
 RL FEBS Lett. 408:5-10(1997).
 CC -1- FUNCTION: CATALYZES THE CLEAVAGE OF L-KYNURENINE AND L-3-
 HYDROXYKYNURENINE INTO ANTHRANILIC AND 3-HYDROXYANTHRANILIC ACIDS,
 CC RESPECTIVELY.
 CC -1- CATALYTIC ACTIVITY: L-KYNURENINE + H(2)O = ANTHRANILATE +
 L-ALANINE.
 CC -1- COFACTOR: PYRIDOXAL PHOSPHATE.
 CC -1- PATHWAY: INVOLVED IN THE BIOSYNTHESIS OF NAD COFACTORS FROM
 TRYPTOPHAN THROUGH THE KYNURENINE PATHWAY.
 CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC.
 CC -1- SIMILARITY: BELONGS TO THE KYNURENINASE FAMILY. SLIGHTLY RELATED
 TO CLASS-V OF PYRIDOXAL-PHOSPHATE-DEPENDENT AMINOTRANSFERASES.
 CC
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 CC
 DR EMBL; U68168; AAC53206.1; --
 DR Hydrolase; Pyridoxal phosphate.
 FT BINDING 276 276 PYRIDOXAL PHOSPHATE (BY SIMILARITY).
 SQ SEQUENCE 464 AA; 52453 MW; 37EE19F0 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 464;
 Best Local Similarity 75.0%; Pred. No. 26;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLAIRLDE 9
 |||||
 Db 28 RVALRLDE 35

RESULT 6
 YWMS_PSEAE ID YWMS_PSEAE STANDARD; PRT; 464 AA.
 AC P28812;
 DT 01-DEC-1992 (Rel. 24, Created)
 DT 01-DEC-1992 (Rel. 24, Last sequence update)
 DT 01-DEC-1992 (Rel. 24, Last annotation update)
 DE HYPOTHETICAL PROTEIN IN MMSB 3'REGION (ORE1) (FRAGMENT).
 OS Pseudomonas aeruginosa.
 OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonas group;
 OC Pseudomonas.
 RN [1]

RP SEQUENCE FROM N.A.
RC STRAIN-ATCC 15692 / PA01;
RX MEDLINE: 92317087.
RA STEELE M.I., LORENZ D., HATTEY K., PARK A., SOKATCH J.R.;
RT "Characterization of the mmsAB operon of Pseudomonas aeruginosa PAO
RT encoding methylmalonate-semialdehyde dehydrogenase and 3-
RT hydroxyisobutyrate dehydrogenase."
RL J. Biol. Chem. 267:13585-13592(1992).
CC -!- SIMILARITY: TO ENZYMES WHICH ACT VIA AN ATP-DEPENDENT COVALENT
CC BINDING OF AMP TO THEIR SUBSTRATE.
CC -----
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CC -----
CC EMBL: M84911; AAA25893.1; -
CC PIR: S27604; S27604.
CC PROSITE: PS00455; AMP-BINDING; 1.
CC PFAM: PF00501; AMP-BINDING; 1.
CC Hypothetical protein.
FT NON_TER 464 464
SQ SEQUENCE 464 AA; 51208 MW; FE491D7C CRC32;

Query Match 65.3%; Score 32; DB 1; Length 464;
Best Local Similarity 55.6%; Pred. No. 26;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9
| | | | |
Db 146 YELALRID 154

RESULT 7
THIS PYRKO
ID THS_PYRKO STANDARD; PRT; 546 AA.
AC Q52500.
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE THERMOSOME SUBUNIT (HEAT-SHOCK PROTEIN).
GN THS.
OS Pyrococcus kodakaraensis.
OC Archaea; Euryarchaeota; Thermococcales; Thermococcaceae; Pyrococcus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-KOD1;
RA IZAWA Y., KAKIHARA H., TAKAGI M., IMANAKA T.;
RL Submitted (APR-1994) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: MOLECULAR CHAPERONE; BINDS UNFOLDED POLYPEPTIDES IN
CC VITRO, AND HAS A WEAK ATPASE ACTIVITY (BY SIMILARITY).
CC -!- SUBUNIT: FORMS AN OLIGOMERIC COMPLEX OF EIGHT-MEMBERED RINGS
CC (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE TCP-1 CHAPERONIN FAMILY.
CC -----
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CC -----
CC EMBL: D29672; BAA06143.1; -
CC HSSP: P48424; IASX.
CC PROSITE: PS00751; TCP1_1; 1.
CC PROSITE: PS00751; TCP1_2; 1.
CC PROSITE: PS00995; TCP1_3; 1.
CC PFAM: PF00118; cpn60_TCP1; 1.

KW Chaperone: ATP-binding; Heat shock.
SQ SEQUENCE 546 AA; 59158 MW; 5B3C9283 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 546;
Best Local Similarity 100.0%; Pred. No. 31;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDE 9
| | | | |
Db 418 LAIRLDE 424

RESULT 8
THSA_THEK1
ID THSA_THEK1 STANDARD; PRT; 548 AA.
AC Q24729;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 15-JUL-1998 (Rel. 36, Last annotation update)
DE THERMOSOME, ALPHA SUBUNIT (CHAPERONIN ALPHA SUBUNIT).
GN THSA.
OS Thermococcus sp. (strain KS-1).
OC Archaea; Euryarchaeota; Thermococcales; Thermococcaceae; Thermococcus.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 98022908.
RA YOSHIDA T., YOHDA M., IIDA T., MARUYAMA T., TAGUCHI H., YAZAKI K.,
RA OHTA T., ODAKA M., ENDO I., KAGAWA Y.;
RT "Structural and functional characterization of homo-oligomeric
RT complexes of alpha and beta chaperonin subunits from the
RT hyperthermophilic archaeum Thermococcus strain KS-1."
RL J. Mol. Biol. 273:635-645(1997).
CC -!- FUNCTION: MOLECULAR CHAPERONE; BINDS UNFOLDED POLYPEPTIDES IN
CC VITRO, AND HAS A WEAK ATPASE ACTIVITY (BY SIMILARITY).
CC -!- SUBUNIT: FORMS AN OLIGOMERIC COMPLEX OF EIGHT-MEMBERED RINGS
CC (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE TCP-1 CHAPERONIN FAMILY.
CC -----
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CC -----
CC EMBL: AB001080; BAA22207.1; -
CC HSSP: P48424; IASX.
CC PROSITE: PS00750; TCP1_1; 1.
CC PROSITE: PS00751; TCP1_2; 1.
CC PROSITE: PS00995; TCP1_3; 1.
CC PFAM: PF00118; cpn60_TCP1; 1.
KW Chaperone: ATP-binding; Multigene family.
SQ SEQUENCE 548 AA; 59191 MW; 08FCFB81 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 548;
Best Local Similarity 100.0%; Pred. No. 32;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDE 9
| | | | |
Db 418 LAIRLDE 424

RESULT 9
GYRA_RICPR
ID GYRA_RICPR STANDARD; PRT; 905 AA.
AC P41080;
DT 01-FEB-1995 (Rel. 31, Created)
DT 01-FEB-1995 (Rel. 31, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)


```
DE DNA GYRASE SUBUNIT A (EC 5.99.1.3).
GN GYRA OR RP206.
OS Rickettsia prowazekii.
OC Bacteria; Proteobacteria; alpha subdivision; Rickettsiales;
OC Rickettsiaceae; Rickettsiae; Rickettsia.
[]]
RN SEQUENCE FROM N.A.
RC STRAIN=MADRID E;
RX MEDLINE; 95129858.
RA WOOD D.O., WAITE R.T.;
RT "Sequence analysis of the Rickettsia prowazekii gyra gene.";
RL Gene 151.191-196(1994).
RN SEQUENCE FROM N.A.
RC STRAIN=MADRID E;
RX MEDLINE; 99039499.
RA ANDERSSON S.G.E., ZOMORODIPOUR A., ANDERSSON J.O.,
RA SICKERITZ-PONTEN T., ALSMARK U.C.M., PODORSKI R.M., NAESLUND A.K.,
RA ERIKSSON A.-S., WINKLER H.H., KURLAND C.G.;
RT "The genome sequence of Rickettsia prowazekii and the origin of
RT mitochondria.";
RL Nature 396:133-140(1998).
CC -|- FUNCTION: DNA GYRASE NEGATIVELY SUPERCOILS CLOSED CIRCULAR DOUBLE-
CC STRANDED DNA IN AN ATP-DEPENDENT MANNER AND ALSO CATALYZES THE
CC INTERCONVERSION OF OTHER TOPOLOGICAL ISOMERS OF DOUBLE-STRANDED
CC DNA RINGS, INCLUDING CATENANES AND KNOTTED RINGS.
CC -|- CATALYTIC ACTIVITY: ATP-DEPENDENT BREAKAGE, PASSAGE AND REJOINING
CC OF DOUBLE-STRANDED DNA.
CC -|- SUBUNIT: MADE UP OF TWO CHAINS. THE A CHAIN IS RESPONSIBLE FOR DNA
CC BREAKAGE AND REJOINING; THE B CHAIN CATALYZES ATP HYDROLYSIS. THE
CC ENZYME FORMS AN A2B2 TETRAMER.
CC
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CC
CC ENBL; U02931; AAA68146.1; -
CC EMBL; AJ235270; CAAL4671.1; -
CC PFAM; PF00521; DNA_topoisomIV; 1.
CC Topoisomerase; Isomerase; DNA-binding.
CC ACT_SITE 123 123 DNA CLEAVAGE (BY SIMILARITY).
CC SEQUENCE 905 AA; 101080 MW; EFBC8ADA CRC32;
CC
CC DR ENBL; U02931; AAA68146.1; -
CC DR EMBL; AJ235270; CAAL4671.1; -
CC DR PFAM; PF00521; DNA_topoisomIV; 1.
CC KW Topoisomerase; Isomerase; DNA-binding.
CC FT ACT_SITE 123 123 DNA CLEAVAGE (BY SIMILARITY).
CC SQ SEQUENCE 905 AA; 101080 MW; EFBC8ADA CRC32;
CC
CC Query Match 65.3%; Score 32; DB 1; Length 905;
CC Best Local Similarity 75.0%; Pred. No. 54;
CC Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
CC
CC QY 2 RLAIRLDE 9
CC :|||||
CC Db 665 KIAIRLDE 672
CC
CC RESULT 10
CC VGLC_HSVE4 STANDARD; PRT; 485 AA.
CC ID VGLC_HSVE4 STANDARD; PRT; 485 AA.
CC AC P22596;
CC DT 01-AUG-1991 (Rel. 19, Created)
CC DT 01-AUG-1991 (Rel. 19, Last sequence update)
CC DT 15-JUL-1999 (Rel. 38, Last annotation update)
CC DE GLYCOPROTEIN.C PRECURSOR (GLYCOPROTEIN 13).
CC GC OR GPI3.
CC GN Equine herpesvirus type 4 (strain 1942) (EHV-4) (Equine herpesvirus
CC OS type 1 subtype 2).
CC OS Viruses; dsDNA viruses, no RNA stage; Herpesviridae;
CC OC Alphaherpesvirinae; Varicellovirus.
CC [1]
CC RN SEQUENCE FROM N.A.
CC RP MEDLINE; 91021040.
```

```
RA NICOLSON L., ONIONS D.E.;
RT "The nucleotide sequence of the equine herpesvirus 4 gc gene
RL homologue.";
RL Virology 179:378-387(1990).
CC -|- SIMILARITY: TO OTHER HERPESVIRUSES GLYCOPROTEIN C.
CC -|- SIMILARITY: BELONGS TO THE IMMUNOGLOBULIN GENE SUPERFAMILY.
CC
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CC
CC ENBL; M58031; AAA46083.1; -
CC EMBL; A21044; CAA01528.1; -
CC PIR; B45343; B45343.
CC KW Glycoprotein; Transmembrane; Signal.
CC FT SIGNAL 1 32
CC FT CHAIN 33 485 GLYCOPROTEIN C.
CC FT DOMAIN 33 444 EXTRACELLULAR.
CC FT TRANSMEM 445 468
CC FT CARBOHYD 60 60 POTENTIAL.
CC FT CARBOHYD 61 61 POTENTIAL.
CC FT CARBOHYD 66 66 POTENTIAL.
CC FT CARBOHYD 67 67 POTENTIAL.
CC FT CARBOHYD 72 72 POTENTIAL.
CC FT CARBOHYD 108 108 POTENTIAL.
CC FT CARBOHYD 116 116 POTENTIAL.
CC FT CARBOHYD 147 147 POTENTIAL.
CC FT CARBOHYD 220 220 POTENTIAL.
CC FT CARBOHYD 225 225 POTENTIAL.
CC FT CARBOHYD 286 286 POTENTIAL.
CC SQ SEQUENCE 485 AA; 52509 MW; 63F72464 CRC32;
CC
CC Query Match 63.3%; Score 31; DB 1; Length 485;
CC Best Local Similarity 60.0%; Pred. No. 45;
CC Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
CC
CC QY 1 YRLAIRLDER 10
CC |||||
CC Db 120 YRLIEHLNQR 129
CC
CC RESULT 11
CC MURF_MYCTU STANDARD; PRT; 510 AA.
CC ID MURF_MYCTU STANDARD; PRT; 510 AA.
CC AC O06220;
CC DT 15-JUL-1998 (Rel. 36, Created)
CC DT 15-JUL-1998 (Rel. 36, Last sequence update)
CC DT 15-JUL-1999 (Rel. 38, Last annotation update)
CC DE UDP-N-ACETYLURAMIDYLALANYL-D-GLUTAMYL-2,6-DIAMINOPIMELATE--D-ALANYL-D-
CC DE ALANYL LIGASE (EC 6.3.2.15) (UDP-MURNAC-PENTAPEPTIDE SYNTHETASE)
CC (D-ALANYL-D-ALANINE-ADDING ENZYME).
CC MURF OR RV2157C OR MTCY270.11.
CC OS Mycobacterium tuberculosis.
CC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; Mycobacterium.
CC Actinomycetales; Corynebacteriaceae; Mycobacteriaceae; Mycobacterium.
CC [1]
CC RN SEQUENCE FROM N.A.
CC RP STRAIN=H37RV;
CC RC MEDLINE; 98295987.
CC RA COLE S.T., BROSCHE R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
CC RA GORDON S.V., EIGLMEIER K., GAS S., BARRY C.E. III, TEKAIA F.,
CC RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
CC RA DAVIES R., DEVLIN K., FELTWEILL T., GENTLES S., HAMLIN N., HOLROYD S.,
CC RA HORNSBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,
CC RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAM M.A., ROGERS J.,
CC RA RUTTER S., SEEGER K., SKELTON S., SQUARES S., SQUARES R., SULSTON J.E.,
CC RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
CC "Deciphering the biology of Mycobacterium tuberculosis from the
```

RT complete genome sequence. ";
RL Nature 393:537-544(1998).
CC -!- FUNCTION: INVOLVED IN CELL WALL FORMATION. CATALYZES THE FINAL
CC STEP IN THE SYNTHESIS OF UDP-N-ACETYLURAMONYL-PENTAPEPTIDE, THE
CC PRECURSOR OF MUREIN (BY SIMILARITY).
CC -!- CATALYTIC ACTIVITY: ATP + UDP-N-ACETYLURAMONYL-L-ALANYL-D-GLUTAMYL
CC -MESO-2,6-DIAMINOHEPTANEDIOATE + D-ALANYL-D-ALANYL = ADP +
CC ORTHOPHOSPHATE + UDP-N-ACETYLURAMONYL-L-ALANYL-D-GAMMA-GLUTAMYL-6-
CC CARBOXY-L-LYSYL-D-ALANYL-D-ALANINE.
CC -!- PATHWAY: PEPTIDOGLYCAN BIOSYNTHESIS.
CC -!- SUBUNIT: MONOMER (BY SIMILARITY).
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC (PROBABLE).
CC -!- SIMILARITY: BELONGS TO THE MURDEF FAMILY.
CC -----
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CC -----
CC EMBL: 295388; CAB08670.1; -
CC PFAM: PF01225; Mur_ligase; 1.
CC Peptidoglycan synthesis; Cell division; Cell wall; Ligase;
KW ATP-binding.
FT NP_BIND 136 142 ATP (POTENTIAL).
SQ SEQUENCE 510 AA; 51632 MW; 4F25A40A CRC32;

Query Match 63.3%; Score 31; DB 1; Length 510;
Best Local Similarity 85.7%; Pred. No. 47;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 2 RLALRLD 8

DB 412 RLALRLD 418

RESULT 12
ESTP DROME
ID ESTP DROME STANDARD; PRT: 544 AA.
AC P18167;
DT 01-NOV-1990 (Rel. 16, Created)
DT 01-NOV-1990 (Rel. 16, Last sequence update)
DT 01-FEB-1996 (Rel. 33, Last annotation update)
DE ESTP DROME P PRECURSOR (EC 3.1.1.1) (EST-P) (CARBOXYLIC-ESTER
DE HYDROLASE).
GN ESTP OR EST-P.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN-CANTON-S;
RX MEDLINE: 90136038.
RA COLLET C., NIELSEN K.M., RUSSELL R.J., KARL M., OAKSHOTT J.G.,
RA RICHMOND R.C.;
RT "Molecular analysis of duplicated esterase genes in Drosophila
RT melanogaster";
RL Mol. Biol. Evol. 7:9-28(1990).
CC -!- CATALYTIC ACTIVITY: A CARBOXYLIC ESTER + H(2)O = AN ALCOHOL
CC + A CARBOXYLIC ANION.
CC -!- SUBUNIT: MONOMER.
CC -!- DEVELOPMENTAL STAGE: MAINLY IN LATE LARVAE.
CC -!- SIMILARITY: BELONGS TO THE TYPE-B CARBOXYLESTERASE/LIPASE FAMILY.
CC -----
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CC -----
CC EMBL: M33780; AAA28520.1; -
CC PIR: B34089; B34089.
CC HSSP: P21836; LMAH.
CC DR FLXBASE; FBG0000594; Est-P.
CC DR PROSITE; PS00122; CARBOXYLESTERASE_B_1; 1.
CC DR PROSITE; PS00941; CARBOXYLESTERASE_B_2; 1.
CC DR PFAM; PF00135; Coesterase; 1.
KW Hydrolase; Serine esterase; Glycoprotein; Signal.
FT SIGNAL 1 19
FT CHAIN 20 544
FT ACT_SITE 206 206
FT ACT_SITE 466 466
FT DISULFID 83 102
FT DISULFID 258 270
FT DISULFID 514 535
FT CARBOHYD 75 75
FT CARBOHYD 114 114
FT CARBOHYD 262 262
FT CARBOHYD 456 456
SQ SEQUENCE 544 AA; 61230 MW; E9F6EEDD CRC32;

Query Match 63.3%; Score 31; DB 1; Length 544;
Best Local Similarity 66.7%; Pred. No. 51;
Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 1 YRLAIRLDE 9

DB 479 YRIGIRPDE 487

RESULT 13
RPC3_YEAST
ID RPC3_YEAST STANDARD; PRT: 654 AA.
AC P32349; Q06591;
DT 01-OCT-1993 (Rel. 27, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE DNA-DIRECTED RNA POLYMERASE III 74 KD POLYPEPTIDE (EC 2.7.7.6) (C74).
GN RPC3 OR RPC82 OR YPR190C OR P9677.11.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomycetes.
RN [1]
RP SEQUENCE FROM N.A. AND PARTIAL SEQUENCE.
RX MEDLINE: 93024385.
RA CHIANNILKULCHAI N., STALDER R., RIVA M., CARLES C., WERNER M.,
RA SENTENAC A.;
RT "RPC82 encodes the highly conserved, third-largest subunit of RNA
RT polymerase C (III) from Saccharomyces cerevisiae";
RL Mol. Cell. Biol. 12:4433-4440(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-S288C / AB972;
RA JOHNSTON M., ANDREWS S., BRINKMAN R., COOPER J., DING H., DU Z.,
RA FAYELLO A., FULTON L., GATUNG S., GRECO T., KIRSTEN J., KUCABA T.,
RA HALLSWORTH K., HAWKINS J., HILLIER L., JIER M., JOHNSON D.,
RA JOHNSTON L., LANGSTON Y., LATREILLE P., LE T., MARDIS E., MENEZES S.,
RA MILLER N., NHAN M., PAULEY A., PELUSO D., RIFKEN L., RILES L.,
RA TATCH A., TREVASKIS E., VIGNATI D., WILCOX L., WOLDMAN P., VAUDIN M.,
RA WILSON R., WATERSTON R.;
RL Submitted (APR-1995) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: DNA-DEPENDENT RNA POLYMERASE CATALYZES THE TRANSCRIPTION
CC OF DNA INTO RNA USING THE FOUR RIBONUCLEOSIDE TRIPHOSPHATES AS
CC SUBSTRATES.
CC -!- CATALYTIC ACTIVITY: N NUCLEOSIDE TRIPHOSPHATE = N PYROPHOSPHATE +
CC RNA(N).
CC -!- SUBUNIT: RNA POLYMERASE III CONSISTS OF ABOUT 15 DIFFERENT
CC SUBUNITS. THIS SUBUNIT IS THE THIRD LARGEST COMPONENT OF RNA
CC POLYMERASE III.

CC -!- SUBCELLULAR LOCATION: NUCLEAR.
CC -!- MISCELLANEOUS: THREE DISTINCT ZINC-CONTAINING RNA POLYMERASES ARE
CC FOUND IN EUKARYOTIC NUCLEI: POLYMERASE I FOR THE RIBOSOMAL RNA
CC PRECURSOR, POLYMERASE II FOR THE MRNA PRECURSOR, AND POLYMERASE
CC III FOR 5S AND TRNA GENES.
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CC
CC EMBL; X63500; CAA45072.1; -
CC DR EMBL; U25841; AAB64619.1; -
CC DR PIR; S31298; S31298.
CC DR SGD; L0001693; RPO82.
CC KW Transferase; DNA-directed RNA polymerase; Transcription; Zinc;
CC Nuclear protein.
CC FT DOMAIN 581 602 LEUCINE-ZIPPER.
CC FT CONFLICT 637 637 V -> L (IN REF. 1).
CC SQ SEQUENCE 654 AA; 74016 MW; 9E17F4F8 CRC32;

Query Match 63.3%; Score 31; DB 1; Length 654;
Best Local Similarity 50.0%; Pred. No. 62;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
|:::|:|:
DB 312 YKIALRLTEQ 321

RESULT 14
YK37_YEAST
ID YK37_YEAST STANDARD; PRT; 705 AA.
AC Q03824;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE HYPOTHETICAL 81.5 KD PROTEIN IN HLJ1-SMP2 INTERGENIC REGION.
GN YMR163C OR YMR520.12C.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomyces.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-S288C / AB972;
RA HUNT S., BOWMAN S., BARRELL B.G., RAJANDREAM M.A.;
RL Submitted (MAY-1995) to the EMBL/GenBank/DDBJ databases.
CC
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CC
CC EMBL; Z49705; CAA89799.1; -
CC DR Hypothetical protein.
CC KW
CC SQ SEQUENCE 705 AA; 81466 MW; 6E07A99F CRC32;

Query Match 63.3%; Score 31; DB 1; Length 705;
Best Local Similarity 55.6%; Pred. No. 67;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9
|:::|:|:
DB 136 YRLSLHLQ 144

RESULT 15
MLB_MYLE
ID MLB_MYLE STANDARD; PRT; 1014 AA.
AC Q06079;
DT 15-DEC-1999 (Rel. 39, Created)
DT 15-DEC-1999 (Rel. 39, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE PUTATIVE MEMBRANE PROTEIN MMPL11.
GN MMPL11 OR MLC1622.16C.
OS Mycobacterium leprae.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RA HAMLIN N., CHURCHER C.M., PARKHILL J., BARRELL B.G., RAJANDREAM M.A.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DDBJ databases.
CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN (POTENTIAL).
CC -!- SIMILARITY: BELONGS TO THE MMPL FAMILY.
CC
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CC
CC EMBL; Z95398; CAB08003.1; -
CC KW Hypothetical protein; Transmembrane.
CC FT TRANSMEM 13 33 POTENTIAL.
CC FT TRANSMEM 188 208 POTENTIAL.
CC FT TRANSMEM 214 234 POTENTIAL.
CC FT TRANSMEM 235 255 POTENTIAL.
CC FT TRANSMEM 279 299 POTENTIAL.
CC FT TRANSMEM 311 331 POTENTIAL.
CC FT TRANSMEM 373 393 POTENTIAL.
CC FT TRANSMEM 409 429 POTENTIAL.
CC FT TRANSMEM 530 550 POTENTIAL.
CC FT TRANSMEM 560 580 POTENTIAL.
CC FT TRANSMEM 598 618 POTENTIAL.
CC FT TRANSMEM 649 669 POTENTIAL.
CC FT TRANSMEM 671 691 POTENTIAL.
CC SQ SEQUENCE 1014 AA; 109875 MW; A2FC256A CRC32;

Query Match 63.3%; Score 31; DB 1; Length 1014;
Best Local Similarity 50.0%; Pred. No. 99;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLDER 10
|:::|:|:
DB 117 YGVSLRLDDR 126

Search completed: February 8, 2000, 00:59:53
Job time: 3782 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:39 ; Search time 209.03 Seconds
(without alignments)
3.317 Million cell updates/sec

Title: US-08-653-294-13
Perfect score: 49
Sequence: 1 YRLAIRLDER 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 225878 seqs, 69334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database :

SPTREMBL_12.*
1: sp_archaea.*
2: sp_bacteria.*
3: sp_fungi.*
4: sp_human.*
5: sp_invertebrate.*
6: sp_mammal.*
7: sp_mhc.*
8: sp_organelle.*
9: sp_phage.*
10: sp_plant.*
11: sp_rodent.*
12: sp_virus.*
13: sp_vertebrate.*
14: sp_unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	38	77.6	1324	2 Q44103	Q44103 amycolatops
2	34	69.4	229	5 Q19415	Q19415 caenorhabdi
3	34	69.4	1154	3 Q01375	Q01375 neurospora
4	34	69.4	1154	3 Q01379	Q01379 neurospora
5	33	67.3	264	2 Q32734	Q32734 agrobacteri
6	33	67.3	554	2 Q06421	Q06421 mycobacteri
7	32	65.3	151	1 Q58407	Q58407 pyrococcy
8	32	65.3	185	2 Q35192	Q35192 synchocyst
9	32	65.3	192	3 Q13610	Q13610 schizosacch
10	32	65.3	304	5 Q9XU51	Q9XU51 caenorhabdi
11	32	65.3	319	6 Q28004	Q28004 bos taurus
12	32	65.3	349	2 Q92H35	Q92H35 enterobacte
13	32	65.3	422	1 Q9WX84	Q9WX84 acidiphiliu
14	32	65.3	548	1 Q9Y813	Q9Y813 pyrococcy
15	32	65.3	1044	3 Q00943	Q00943 pichia angu
16	31	63.3	236	2 Q9X0G1	Q9X0G1 thermotoga
17	31	63.3	246	2 Q54045	Q54045 pseudomonas
18	31	63.3	277	2 Q07463	Q07463 rhodospseudo
19	31	63.3	302	5 Q25608	Q25608 onchocerca
20	31	63.3	356	10 Q9XFP1	Q9XFP1 arabidopsis

21	31	63.3	456	5 Q9XX98	Q9XX98 caenorhabdi
22	31	63.3	463	1 Q9YAG3	Q9YAG3 aeropyrum p
23	31	63.3	485	12 Q39258	Q39258 equine herp
24	31	63.3	517	2 Q69556	Q69556 mycobacteri
25	31	63.3	573	4 Q94830	Q94830 homo sapien
26	31	63.3	740	10 Q82436	Q82436 cucumis mel
27	31	63.3	741	10 Q81122	Q81122 malus domes
28	31	63.3	845	5 Q01914	Q01914 caenorhabdi
29	31	63.3	1110	3 Q92198	Q92198 aspergillus
30	31	63.3	1254	13 Q9YHU2	Q9YHU2 brachydanio
31	31	63.3	1524	1 Q54436	Q54436 staphylothe
32	31	63.3	1699	5 Q44344	Q44344 strongyloce
33	31	63.3	2282	11 Q61479	Q61479 mus musculu
34	30	61.2	35	12 Q65737	Q65737 bluetongue
35	30	61.2	124	2 Q05227	Q05227 bacillus su
36	30	61.2	125	1 Q9YBF7	Q9YBF7 aeropyrum p
37	30	61.2	128	2 Q33420	Q33420 pseudomonas
38	30	61.2	130	2 Q05281	Q05281 escherichia
39	30	61.2	135	2 Q52150	Q52150 escherichia
40	30	61.2	135	2 Q85624	Q85624 escherichia
41	30	61.2	141	2 Q67689	Q67689 aquifex aso
42	30	61.2	199	5 Q27010	Q27010 toxoplasma
43	30	61.2	284	5 Q23366	Q23366 caenorhabdi
44	30	61.2	312	5 Q17557	Q17557 caenorhabdi
45	30	61.2	318	5 Q44027	Q44027 toxoplasma

ALIGNMENTS

RESULT 1

Q44103 ID Q44103 PRELIMINARY; PRG: 1324 AA.
AC Q44103;
DT 01-NOV-1996 (TREMREL. 01, Created)
DT 01-NOV-1996 (TREMREL. 01, Last sequence update)
DT 01-NOV-1999 (TREMREL. 12, Last annotation update)
DE PEPTIDE-SYNTHETASE (FRAGMENT).
GN APS.

OS Amycolatopsis mediterranei.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae; Amycolatopsi.
OC Actinomycetales; Pseudonocardineae; Pseudonocardaceae; Amycolatopsi.
RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=DSM 5908;

RX MEDLINE: 97449857

RA PELZER S., REICHERT W., HUPPERT M., HECKMANN D., WOHLLEBEN W.;

RT "Cloning and analysis of a peptide synthetase gene of the baohimycin producer Amycolatopsis mediterranei DSM5908 and development of a gene disruption/replacement system.";

RL J. Biotechnol. 56:115-128(1997).

DR EMBL: X97860; CAA66454.1; -

DR PROSITE; PS00455; AMP_BINDING; 1.

DR PFAM; PF00501; AMP-binding; 1.

DR PFAM; PF00668; DUF4; 2.

KW Ligase.

FT NON_TER 1 1324

FT NON_TER 1324 1324

SQ SEQUENCE 1324 AA; 142666 MW; 2C08588E CRC32;

Query Match 77.6%; Score 38; DB 2; Length 1324;

Best Local Similarity 80.08; Pred. No. 20;

Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Oy 1 YRLAIRLDER 10

Db 969 YRVAGRLDER 978

RESULT 2

Q19415

ID Q19415 PRELIMINARY; PRT; 229 AA.
 AC Q19415;
 DT 01-NOV-1996 (TREMELREL. 01, Created)
 DT 01-NOV-1996 (TREMELREL. 01, Last sequence update)
 DT 01-JAN-1999 (TREMELREL. 09, Last annotation update)
 DE F13E9.10 PROTEIN.
 GN F13E9.10.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MCMURRAY A.;
 RL Submitted (FEB-1996) to the EMBL/GenBank/DBJ databases.
 [2]
 RP SEQUENCE FROM N.A.
 RN [1]
 RX MEDLINE; 94150718.
 RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,
 RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
 RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
 RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,
 RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
 RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
 RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
 RA SWALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
 RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
 RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 RT elegans.";
 RL Nature 368:32-38(1994).
 DR EMBL; 269383; CAA93413.1; -.
 SQ SEQUENCE 229 AA; 26620 MW; F822FE98 CRC32;

Query Match 59.4%; Score 34; DB 5; Length 229;
 Best Local Similarity 70.0%; Pred. No. 21;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 1 YRLAIRLDR 10
 Db 115 YEQAIRLDR 124

RESULT 3
 QO1375 PRELIMINARY; PRT; 1154 AA.
 ID QO1375
 AC QO1375;
 DT 01-NOV-1996 (TREMELREL. 01, Created)
 DT 01-NOV-1996 (TREMELREL. 01, Last sequence update)
 DT 01-NOV-1998 (TREMELREL. 08, Last annotation update)
 DE HYPOTHETICAL 130.4 KD PROTEIN.
 OS Neurospora crassa.
 OC Eukaryota; Fungi; Ascomycota; Euascomycetes; Pyrenomycetes;
 OC Sordariales; Sordariaceae; Neurospora.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-J1518;
 RX MEDLINE; 94203179.
 RA CAMBARERI E.B., HELBER J., KINSEY J.A.;
 RT "Rad1-1, an active LINE-like element of Neurospora crassa.";
 RL Mol. Gen. Genet. 242:658-665(1994).
 DR EMBL; L25662; AAA21781.1; -.
 DR PFAM; PF00078; rvt; 1.
 KW Hypothetical protein.
 FT DOMAIN 1019 1022 POLY-LYS.
 FT DOMAIN 1029 1034 POLY-GLU.
 SQ SEQUENCE 1154 AA; 130398 MW; DF0BA680 CRC32;

Query Match 59.4%; Score 34; DB 3; Length 1154;
 Best Local Similarity 66.7%; Pred. No. 40;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 RLAIRLDR 10
 Db 246 RLAVKLDR 254

OY 1 YRLAIRLDR 9
 Db 1136 YRLAVELEE 1144

RESULT 4
 QO1379 PRELIMINARY; PRT; 1154 AA.
 ID QO1379
 AC QO1379;
 DT 01-NOV-1996 (TREMELREL. 01, Created)
 DT 01-NOV-1996 (TREMELREL. 01, Last sequence update)
 DT 01-NOV-1998 (TREMELREL. 08, Last annotation update)
 DE HYPOTHETICAL 130.5 KD PROTEIN.
 OS Neurospora crassa.
 OC Eukaryota; Fungi; Ascomycota; Euascomycetes; Pyrenomycetes;
 OC Sordariales; Sordariaceae; Neurospora.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-J1518;
 RX MEDLINE; 94203179.
 RA CAMBARERI E.B., HELBER J., KINSEY J.A.;
 RT "Rad1-1, an active LINE-like element of Neurospora crassa.";
 RL Mol. Gen. Genet. 242:658-665(1994).
 DR EMBL; L25663; AAA21792.1; -.
 DR PFAM; PF00078; rvt; 1.
 KW Hypothetical protein.
 FT DOMAIN 1019 1022 POLY-LYS.
 FT DOMAIN 1029 1034 POLY-GLU.
 SQ SEQUENCE 1154 AA; 130470 MW; 7FBE8EAF CRC32;

Query Match 69.4%; Score 34; DB 3; Length 1154;
 Best Local Similarity 66.7%; Pred. No. 1.2e+02;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 1 YRLAIRLDR 9
 Db 1136 YRLAVELEE 1144

RESULT 5
 O32734 PRELIMINARY; PRT; 264 AA.
 ID O32734
 AC O32734;
 DT 01-JAN-1998 (TREMELREL. 05, Created)
 DT 01-JAN-1998 (TREMELREL. 05, Last sequence update)
 DT 01-NOV-1999 (TREMELREL. 12, Last annotation update)
 DE AITE PROTEIN.
 GN AITE.
 OS Agrobacterium tumefaciens.
 OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
 OC Rhizobiaceae; Agrobacterium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-C58;
 RX MEDLINE; 96359388.
 RA MATTHYSSE A.G., VARNALL H.A., YOUNG N.;
 RT "Requirement for genes with homology to ABC transport systems for
 RT attachment and virulence of Agrobacterium tumefaciens.";
 RL J. Bacteriol. 178:5302-5308(1996).
 DR EMBL; U59485; AAB67299.1; -.
 DR PFAM; PF00005; ABC tran; 1.
 SQ SEQUENCE 264 AA; 28745 MW; D5629761 CRC32;

Query Match 67.3%; Score 33; DB 2; Length 264;
 Best Local Similarity 66.7%; Pred. No. 40;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 RLAIRLDR 10
 Db 246 RLAVKLDR 254

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RESULT 6
O06421 PRELIMINARY; PRT; 554 AA.
AC O06421;
DT 01-JUL-1997 (TReMBLrel. 04, Created)
DT 01-JUL-1997 (TReMBLrel. 04, Last sequence update)
DT 01-NOV-1998 (TReMBLrel. 08, Last annotation update)
DE MEND.
DE MEND.
GN MEND.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-H37RV;
RA BROWN D., CHURCHER C.M.;
RA Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-H37RV;
RA BARRELL B.G., RAJANDREAM M.A.;
RA Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-H37RV;
RA MEDLINE; 96181548.
RA PHILIPP W.J., POULET S., EIGLMEIER K., PASCOPELLA L.,
RA BALASUBRAMANIAN V., HEYM B., BERGH S., BLOOM B.R., JACOBS W.R. JR.,
RA COLE S.T.;
RT "An integrated map of the genome of the tubercle bacillus,
RT Mycobacterium tuberculosis H37RV, and comparison with Mycobacterium
RT leprae.";
RL Proc. Natl. Acad. Sci. U.S.A. 93:3132-3137(1996).
DR EMBL: Z95558; CAB08966.1; -.
SQ SEQUENCE 554 AA; 57835 MW; C42C89FC CRC32;

Query Match 67.3%; Score 33; DB 2; Length 554;
Best Local Similarity 66.7%; Pred. No. 87;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLAI RLDR 10
Db 48 RLVRIDER 56

RESULT 7
O58407 PRELIMINARY; PRT; 151 AA.
AC O58407;
DT 01-AUG-1998 (TReMBLrel. 07, Created)
DT 01-AUG-1998 (TReMBLrel. 07, Last sequence update)
DT 01-JAN-1999 (TReMBLrel. 09, Last annotation update)
DE 151AA LONG HYPOTHETICAL FRAX PROTEIN.
GN PH0674.
OS Pyrococcus horikoshii.
OC Archaea; Euryarchaeota; Thermococcales; Thermococcaceae; Pyrococcus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-OT3;
RX MEDLINE; 98344137.
RA KAWARABAYASI Y., SAWADA M., HORIKAWA H., HAIKAWA Y., HINO Y.,
RA YAMAMOTO S., SEKINE M., BABA S., KOSUGI H., HOSIYAMA A., NAGAI Y.,
RA SAKAI M., OGURA K., OTUKA R., NAKAZAWA H., TAKAMIYA M., OHFUKU Y.,
RA FUNAHASHI T., TANAKA T., KUDOH Y., YAMAZAKI J., KUSHIDA N., OGUCHI A.,
RA AOKI K., NAKAMURA Y., ROBB T.F., HORIKOSHI K., MASUCHI Y., SHIZUYA H.,
RA KIKUCHI H.;
RT "Complete sequence and gene organization of the genome of a hyper-
RT thermophilic archaeobacterium, Pyrococcus horikoshii OT3.";
RL DNA Res. 5:55-76(1998).
DR EMBL: AP000003; BAA29765.1; -.
SQ SEQUENCE 151 AA; 17160 MW; 11AACD59 CRC32;
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Query Match 65.3%; Score 32; DB 1; Length 151;
Best Local Similarity 77.8%; Pred. No. 36;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 2 RLAI RLDR 10
Db 96 RLII RLDR 104

RESULT 8
O55192 PRELIMINARY; PRT; 185 AA.
ID O55192;
AC O55192;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
DT 01-JAN-1999 (TReMBLrel. 09, Last annotation update)
DE HYPOTHETICAL 20.8 KD PROTEIN.
OS Synechocystis sp. (strain PCC 6803).
OC Bacteria; Cyanobacteria; Chroococcales; Synechocystis.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-PCC6803;
RA TABATA S.;
RA Submitted (AUG-1995) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-PCC6803;
RX MEDLINE; 96127529.
RA KANEKO T., TANAKA A., SATO S., KOTANI H., SAZUKA T., MIYAJIMA N.,
RA SUGIURA M., TABATA S.;
RT "Sequence analysis of the genome of the unicellular cyanobacterium
RT Synechocystis sp. strain PCC6803. I. Sequence features in the 1 Mb
RT region from map positions 64% to 92% of the genome.";
RL DNA Res. 2:153-166(1995).
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN-PCC6803;
RX MEDLINE; 97061201.
RA KANEKO T., SATO S., KOTANI H., TANAKA A., ASAMIZU E., NAKAMURA Y.,
RA MIYAJIMA N., HIROSAWA M., SUGIURA M., SASAMOTO S., KIMURA T.,
RA HOSOUCHI T., MATSUNO A., MURAKI A., NAKAZAKI N., NARUO K., OKUMURA S.,
RA SHIMPO S., TAKEUCHI C., WADA T., WATANABE A., YAMADA M., YASUDA M.,
RA TABATA S.;
RT "Sequence analysis of the genome of the unicellular cyanobacterium
RT Synechocystis sp. strain PCC6803. II. Sequence determination of the
RT entire genome and assignment of potential protein-coding regions.";
RL DNA Res. 3:109-136(1996).
DR EMBL: D64001; BAA10334.1; -.
KW Hypothetical protein.
SQ SEQUENCE 185 AA; 20830 MW; 365A078D CRC32;

Query Match 65.3%; Score 32; DB 2; Length 185;
Best Local Similarity 60.0%; Pred. No. 44;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 1 YRLAI RLDR 10
Db 35 YRLALRIQLR 44

RESULT 9
O13610 PRELIMINARY; PRT; 192 AA.
ID O13610;
AC O13610;
DT 01-JAN-1998 (TReMBLrel. 05, Created)
DT 01-JAN-1998 (TReMBLrel. 05, Last sequence update)
DT 01-AUG-1998 (TReMBLrel. 07, Last annotation update)
DE HYPOTHETICAL 22.4 KD PROTEIN.
GN P1019.
OS Schizosaccharomyces pombe (fission yeast).
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OC Eukaryota; Fungi; Ascomycota; Archiascomycetes;
OC Schizosaccharomycetales; Schizosaccharomycetaceae;
OC Schizosaccharomycetes.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-972 H-;
RA KUSHIDA N., YAMAZAKI S., TANAKA T., JINNO K., HAIKAWA Y., YAMAZAKI J.,
RA YAMAMOTO S., SEKINE M., OGUCHI A., NAGAI Y., SAKAI M., AOKI K.,
RA OCURA K., OTSUKA R., KUDOH Y., YANAGIDA M., MACHIDA M., ZHANG M.Q.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL: AS004535; BAA21398.1;
KW Hypothetical protein
SQ SEQUENCE 192 AA; 22352 MW; F43F0759 CRC32;

Query Match 65.3%; Score 32; DB 3; Length 192;
Best Local Similarity 58.3%; Pred. No. 46;
Matches 7; Conservative 3; Mismatches 0; Indels 2; Gaps 1;

QY 1 YRLAIRL--DER 10
Db 115 YRLAIRAKDER 126
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||:|:| |

RESULT 10
Q9XU51
ID Q9XU51 PRELIMINARY; PRT; 304 AA.
AC Q9XU51
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE T08G3.6 PROTEIN.
GN T08G3.6.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA LLOYD C.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans";
RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA WILSON R., AINSKOUGH R., ANDERSON K., BAYNES C., BERKS M.,
RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
RA SHALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans";
RL Nature 368:32-38(1994).
DR EMBL; 283238; CAB05796.1;
SQ SEQUENCE 304 AA; 34207 MW; F9701C2D CRC32;

Query Match 65.3%; Score 32; DB 5; Length 304;
Best Local Similarity 77.8%; Pred. No. 75;
Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLAIRLDE 9
Db 287 YRLAIRNDE 295
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||||| |

RESULT 11
Q9XU51
ID Q9XU51 PRELIMINARY; PRT; 304 AA.
AC Q9XU51
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE T08G3.6 PROTEIN.
GN T08G3.6.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA LLOYD C.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans";
RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA WILSON R., AINSKOUGH R., ANDERSON K., BAYNES C., BERKS M.,
RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
RA SHALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
RT elegans";
RL Nature 368:32-38(1994).
DR EMBL; 283238; CAB05796.1;
SQ SEQUENCE 304 AA; 34207 MW; F9701C2D CRC32;

Query Match 65.3%; Score 32; DB 2; Length 349;
Best Local Similarity 75.0%; Pred. No. 86;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLD 8
Db 277 YRLAIRID 284
||||| |
||||| |

RESULT 13
Q9XU51
ID Q9XU51 PRELIMINARY; PRT; 422 AA.
AC Q9XU51
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)

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Q28004
ID Q28004 PRELIMINARY; PRT; 319 AA.
AC Q28004;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
DE 11-CIS-RETINOL DEHYDROGENASE.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovinae; Bos.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 95386398.
RA DRIESSEN C.A., JANSSEN B.P., WINKENS H.J., VAN VUGT A.H., LESUW T.L.,
RA JANSSEN J.J.;
RT "Cloning and expression of a cDNA encoding bovine retinal pigment
RT epithelial 11-cis retinol dehydrogenase";
RL Invest. Ophthalmol. Vis. Sci. 36:1988-1996(1995).
DR EMBL; L36533; AAA80694.1;
DR HSSP; P14061; 1FDM.
DR PFAM; PF00106; adh_short; 1.
DR PRINTS; PRO0080; ALCDHDSGNASE.
SQ SEQUENCE 319 AA; 34400 MW; 37A78DAA CRC32;

Query Match 65.3%; Score 32; DB 6; Length 319;
Best Local Similarity 75.0%; Pred. No. 79;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDER 10
Db 45 LAIRLDER 52
||:|:| |
||:|:| |

RESULT 12
Q92H35
ID Q92H35 PRELIMINARY; PRT; 349 AA.
AC Q92H35;
DT 01-MAY-1999 (TREMBlrel. 10, Created)
DT 01-MAY-1999 (TREMBlrel. 10, Last sequence update)
DT 01-MAY-1999 (TREMBlrel. 10, Last annotation update)
DE NITROGEN REGULATORY PROTEIN.
GN NTRB.
OS Enterobacter gergoviae.
OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
OC Enterobacter.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-57-7;
RA DONG Y.M., LI J.D.;
RT "The cloning of glnA, ntrB, and ntrC from Enterobacter gergoviae 57-7
RT and their characterization";
RL Submitted (OCT-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF072440; AAC69321.1;
SQ SEQUENCE 349 AA; 38412 MW; A9F4BA43 CRC32;

Query Match 65.3%; Score 32; DB 2; Length 349;
Best Local Similarity 75.0%; Pred. No. 86;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRLD 8
Db 277 YRLAIRID 284
||||| |
||||| |

RESULT 13
Q9XU51
ID Q9XU51 PRELIMINARY; PRT; 422 AA.
AC Q9XU51
DT 01-NOV-1999 (TREMBlrel. 12, Created)
DT 01-NOV-1999 (TREMBlrel. 12, Last sequence update)

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DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
 DE LIGHT-INDEPENDENT PROTOCHLOROPHYLLIDE REDUCTASE.
 GN BCNN.
 OS Acidiphilium rubrum.
 OC Bacteria: Proteobacteria; alpha subdivision; Acetobacteraceae;
 OC Acidiphilium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MASUDA T., INOUE K., MASUDA M., NAGAYAMA M., OHTA H., SHIMADA H.,
 RA TAKAMURA K.
 RT "The metal-insertion step of bacteriochlorophyll biosynthesis in an
 RT aerobic bacterium Acidiphilium rubrum, which produces zinc-containing
 RT bacteriochlorophyll as natural photosynthetic pigment."
 RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AB017351; BAA76536.1; -.
 SQ SEQUENCE 422 AA; 45864 MW; 4248EA89 CRC32;

Query Match 65.3%; Score 32; DB 2; Length 422;
 Best Local Similarity 77.8%; Pred. No. 1.1e+02;
 Matches 7; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 LAIRLDER 10
 Db 123 RAAALDER 131

RESULT 14
 QY8I3 PRELIMINARY; PRT; 548 AA.
 AC QY8I3;
 DT 01-NOV-1999 (TReMBLrel. 12, Created)
 DT 01-NOV-1999 (TReMBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
 DE CHAPERONIN LIKE PROTEIN ALPHA SUBUNIT.
 GN CPKA.
 OS Pyrococcus kodakaraensis.
 OC Archaea; Euryarchaeota; Thermococcales; Thermococcaceae; Pyrococcus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-KOD1.
 RX MEDLINE; 99203147.
 RA IZUMI M., FUJIWARA S., TAKAGI M., KANAYA S., IMANAKA T.;
 RT "Isolation and characterization of a second subunit of molecular
 RT chaperonin from pyrococcus kodakaraensis KOD1: analysis of an ATPase-
 RT deficient mutant enzyme."
 RL Appl. Environ. Microbiol. 65:1801-1805(1999).
 DR EMBL; AB018432; BAA76952.1; -.
 DR PROSITE; PS00750; TCPL_1; 1.
 DR PROSITE; PS00751; TCPL_2; 1.
 DR PROSITE; PS00995; TCPL_3; 1.
 SQ SEQUENCE 548 AA; 59169 MW; 10DA6C62 CRC32;

Query Match 65.3%; Score 32; DB 1; Length 548;
 Best Local Similarity 100.0%; Pred. No. 1.4e+02;
 Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDE 9
 Db 418 LAIRLDE 424

RESULT 15
 QY0943 PRELIMINARY; PRT; 1044 AA.
 AC QY0943;
 DT 01-NOV-1996 (TReMBLrel. 01, Created)
 DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
 DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
 DE NITRITE REDUCTASE.
 GN YN11.
 OS Pichia angusta (Yeast) (Hansenula polymorpha).

OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Pichia.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-NCYC 495;
 RA BRITO N., AVILA J., PEREZ M., GONZALEZ C., SIVERIO J.M.;
 RL J. Biochem. 317:89-95(1996).
 DR EMBL; Z68122; CAA92206.1; -.
 DR PFAM; PF01077; NIR_SIR; 1.
 DR PFAM; PF00355; Rieske; 1.
 DR PRINTS; PR00397; SIROHAEM.
 SQ SEQUENCE 1044 AA; 116574 MW; 59F4D4B1 CRC32;

Query Match 65.3%; Score 32; DB 3; Length 1044;
 Best Local Similarity 75.0%; Pred. No. 2.7e+02;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 3 LAIRLDER 10
 Db 696 LAVRLEER 703

Search completed: February 8, 2000, 13:17:41
 Job time: 32490 sec

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OM of: US-08-653-294-13 to: GenEmbl.* out_format : pfs
 Date: Feb 8, 2000 4:39 PM
 About: Results were produced by the GenCore software, version 4.5,
 Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:
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 -O=/cgnl1/USPTO.spool/US08653294/runat_04022000_160701_15779/app_query.fasta.1
 -D=GenEmbl -QEMT=fastap -SUFFIX=rge -GAPOP=12.000 -GAPEXT=4.000
 -MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000 -CGAPOP=4.500
 -CGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
 -FGAPEXT=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DELOP=6.000
 -DLEXT=7.000 -START=1 -MATRIX=biosum62 -TRANS=human40.cdi
 -LIST=45 -DLOCALIGN=200 -THR SCORE=pct -ALIGN=15 -MODE=LOCAL
 -OUTFWT=pfs -NORM=ext -MINLEN=0 -MAXLEN=1000000 -USER=US08653294
 -NCPU=6 -ICPU=3 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-08-653-294-13
 Query length: 10
 Database: GenEmbl.*
 Database sequences: 821193
 Database length: -1518192014
 Search time (sec): 11370.480000

score_list:

Sequence	Strd Orig	ZScore	Escore	Len	Documentation
gb_pr2:HS1106N18	-	40.00	96.64	2.3e+03	142336 ! AL035457 Human DNA sequence
gb_pr2:AL004965	-	40.00	89.16	6.1e+03	323792 ! AC004965 Homo sapiens clone
gb_cm:AF064209	-	39.00	126.37	51.38	3452 ! AF064209 Macropus eugenii inte
gb_in2:AC008207	-	39.00	103.02	1.0e+03	44887 ! AC018207 Drosophila melanogas
gb_in2:AC005268	-	39.00	96.72	2.3e+03	89731 ! AC005268 Drosophila melanogas
gb_ba1:AF064209	+	38.00	120.97	102.66	3975 ! X97860 Amycolatopsis mediterr
gb_pr2:HS204ES	+	38.00	89.24	6.0e+03	129969 ! X98941 Human DNA sequence fr
gb_pr4:AC009239	+	38.00	88.80	6.4e+03	136371 ! AC009239 Homo sapiens clone
gb_htg4:AC001130	+	38.00	86.97	8.0e+03	166832 ! AC011830 Homo sapiens chrom
gb_htg3:AC010086	+	38.00	85.91	9.2e+03	187447 ! AC010086 Homo sapiens clone
gb_htg4:AC008591	+	38.00	84.80	1.1e+04	211769 ! AC008591 Homo sapiens chrom
gb_in1:AF026152	+	37.00	135.98	14.97	486 ! AF026152 Caenorhabditis elegans
gb_ba1:HSW08XMXU	+	37.00	126.87	48.18	1323 ! Y09104 Hyphomicrobium sp. DNA,
gb_in1:CELC04F6	+	37.00	100.09	1.5e+03	25083 ! U42835 Caenorhabditis elegans
gb_in1:CELF09F7	+	37.00	99.72	1.6e+03	26121 ! AC015346 Drosophila melanogas
gb_in1:CELF09F7	+	37.00	97.82	2.0e+03	32202 ! U00050 Caenorhabditis elegans
gb_htg6:AC011071	-	37.00	84.34	1.1e+04	141615 ! AC011071 Drosophila melanog
gb_htg4:AC008274	+	37.00	83.49	1.3e+04	155224 ! AC008274 Homo sapiens clone
gb_in2:AC002465	+	37.00	83.47	1.3e+04	155881 ! AC002465 Human BAC clone RG3
gb_htg6:AC010847	-	37.00	82.44	1.4e+04	174541 ! AC010847 Drosophila melanog
gb_htg3:AC008537	-	37.00	81.80	1.6e+04	187246 ! AC008537 Homo sapiens chrom
gb_pr2:HSW800934	-	36.00	112.35	310.04	4150 ! AL117426 Homo sapiens mRNA: cl
gb_in1:DMUSHAPE	-	36.00	111.11	363.76	4759 ! Y12322 D.melanogaster mRNA for
gb_in1:CEW08G11	-	36.00	94.76	3.0e+03	28690 ! X28817 Caenorhabditis elegans
gb_htg3:AC010400	-	36.00	92.45	4.0e+03	36974 ! AC010400 Homo sapiens chromos
gb_in2:AC004368	-	36.00	84.37	1.1e+04	98985 ! AC004368 Drosophila melanog
gb_in2:AC005269	-	36.00	83.02	1.3e+04	104378 ! AC005269 Drosophila melanog
gb_pr3:HS281H8	+	36.00	82.50	1.4e+04	110414 ! AC031133 Human DNA sequence
gb_htg1:AP000585	+	36.00	81.48	1.6e+04	123497 ! AP000585 Homo sapiens chrom
gb_htg6:AC008002	+	36.00	81.25	1.7e+04	126629 ! AC008002 Drosophila melanog
gb_htg7:AC001715	+	36.00	78.90	2.2e+04	163954 ! AC011755 Drosophila melanog
gb_htg1:AP000642	+	36.00	78.56	2.3e+04	170145 ! AP000642 Homo sapiens chrom
gb_ov:AF073712	+	35.00	125.79	55.32	603 ! AF073712 Salvelinus fontinalis
gb_pr2:HS01BRR	+	35.00	124.62	64.30	686 ! AL114607 Botrytis cinerea strai
gb_p12:CN501AKG	+	35.00	123.45	74.70	780 ! AL112548 Botrytis cinerea strai
gb_p11:CN501AKG	+	35.00	119.56	123.01	1196 ! X55548 A.nidulans gene for con
gb_pr2:HS011741	+	35.00	114.45	236.75	2096 ! AJ011741 Homo sapiens TOP2 a
gb_p11:ENU12630	-	35.00	112.21	315.82	2683 ! U12630 Emeritella nidulans rib
gb_in2:CEU58085	-	35.00	111.15	361.60	3013 ! U58085 Caenorhabditis elegans
gb_p11:SCLPV1	+	35.00	109.43	450.72	3639 ! X67315 S.cerevisiae Lxp1 gene
gb_ba2:AE000541	+	35.00	97.84	2.0e+03	13012 ! AE000541 Helicobacter pylori
gb_htg3:AC014153	+	35.00	90.49	5.1e+03	29183 ! AC014153 Drosophila melanogas

gb_pr3:AC005605 - 35.00 88.18 6.9e+03 37640 ! AC005605 Homo sapiens subt
 gb_p11:AB015475 + 35.00 80.84 1.8e+04 84325 ! AB015475 Arabidopsis thali
 gb_p12:ATF1715 - 35.00 80.58 1.8e+04 86748 ! AL031032 Arabidopsis thali

seq_name: gb_pr2:HS1106N18

seq_documentation_block:

LOCUS HS1106N18 142336 bp DNA PRI 23-NOV-1999
 DEFINITION Human DNA sequence from clone 1106N18 on chromosome 20q13.2-13.2,
 complete sequence.

ACCESSION AL035457

VERSION AL035457.13 GI:6143575

KEYWORDS HTG.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (Bases 1 to 142336)

AUTHORS Clark,G.

TITLE Direct Submission

JOURNAL Submitted (30-OCT-1999) Sanger Centre, Hinxton, Cambridgeshire,
 CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

requests: clonerequest@sanger.ac.uk

On Oct 29, 1999 this sequence version replaced gi:6065875.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

The following abbreviations are used to associate primary accession

numbers given in the feature table with their source databases:

En: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information

on the WORMPEP database can be found at

http://www.sanger.ac.uk/projects/C.elegans/wormep/important: This

sequence is not the entire insert of clone 1106N18. It may be

shorter because we only sequence overlapping sections once, or

longer because we arrange for a small overlap between neighbouring

submissions.

The true left end of clone 906P16 is at 142237 in this sequence.

The true right end of clone dj1193N1 is at 79367 in this sequence.

This sequence has been finished according to sequence map criteria

as follows. An attempt is made to resolve all sequencing problems,

such as compressions and repeats, but not necessarily within known

annotated human repeat sequence elements (e.g. Alu). Where the

sequence is ambiguous, there is an annotation using the 'unsure'

feature key.

This sequence was generated from part of bacterial clone contigs of

human chromosome 20, constructed by the Sanger Centre Chromosome 20

Mapping Group. Further information can be found at

http://www.sanger.ac.uk/HGP/Chr20

1106N18 is from the library RPCI-5 constructed at the Roswell Park

Cancer Institute by the group of Pieter de Jong. For further

details see http://bacpac.med.buffalo.edu/VECTOR: PCYAC2.

Location/Qualifiers

1. 142336

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="20"

/map="q13.2-13.2"

/clone_lib="RPCI-5"

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BASE COUNT 37456 a 31918 c 32886 g 40076 t

ORIGIN

alignment_scores:

Quality: 40.00 Length: 10

Ratio: 4.000 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-13 x HS1106N18/rev ..

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*	1849	1867:	gap of unknown length
*	1868	4168:	contig of 2301 bp in length
*	4169	4187:	gap of unknown length
*	4188	6023:	contig of 1836 bp in length
*	6024	6042:	gap of unknown length
*	6043	7886:	contig of 1844 bp in length
*	7887	7905:	gap of unknown length
*	7906	9644:	contig of 1739 bp in length
*	9645	9663:	gap of unknown length
*	9664	11563:	contig of 1900 bp in length
*	11564	11582:	gap of unknown length
*	11583	13437:	contig of 1855 bp in length
*	13438	13456:	gap of unknown length
*	13457	15087:	contig of 1631 bp in length
*	15088	15106:	gap of unknown length
*	15107	17659:	contig of 2553 bp in length
*	17660	17678:	gap of unknown length
*	17679	19643:	contig of 1965 bp in length
*	19644	19662:	gap of unknown length
*	19663	21474:	contig of 1812 bp in length
*	21475	21493:	gap of unknown length
*	21494	23992:	contig of 2499 bp in length
*	23993	24011:	gap of unknown length
*	24012	27269:	contig of 3258 bp in length
*	27270	27288:	gap of unknown length
*	27289	30064:	contig of 2776 bp in length
*	30065	30083:	gap of unknown length
*	30084	33836:	contig of 3753 bp in length
*	33837	33855:	gap of unknown length
*	33856	36884:	contig of 2829 bp in length
*	36885	36703:	gap of unknown length
*	36704	39501:	contig of 2798 bp in length
*	39502	39520:	gap of unknown length
*	39521	43003:	contig of 3483 bp in length

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seq_documentation_block:
LOCUS       AF064209      3452 bp      DNA          MAM          05-OCT-1999
DEFINITION   Macropus eugenii interleukin-5 (IL5) gene, complete cds.
ACCESSION   AF064209
VERSION     AF064209.1 GI:5006325
KEYWORDS
SOURCE      tammar wallaby.
ORGANISM    Macropus eugenii
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
            Metatheria; Diprotodontia; Macropodidae; Macropus.
REFERENCE   1 (bases 1 to 3452)
AUTHORS     Hawken,R.J., Maccarone,P., Toder,R., Marshall Graves,J.A. and
            Maddox,J.F.
TITLE       Isolation and characterization of marsupial IL5 genes
JOURNAL     Immunogenetics 49 (11-12), 942-948 (1999)
MEDLINE     99432005
REFERENCE   2 (bases 1 to 3452)
AUTHORS     Hawken,R.J., Maccarone,P., Toder,R., Marshall Graves,J.A. and
            Maddox,J.F.
TITLE       Direct Submission
JOURNAL     Submitted (07-MAY-1998) Veterinary Pathobiology, University of
            Minnesota, 1988 Fitch Avenue, St. Paul, MN 55108, USA
FEATURES
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BASE COUNT  1046 a 585 c 616 g 1205 t
ORIGIN
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Quality: 39.00 Length: 10
Ratio: 4.333 Gaps: 0
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alignment_block:
US-08-653-294-13 x AF064209/rev ..
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seq_name: gb_htg7:AC018207
seq_documentation_block:
LOCUS       AC018207      44887 bp      DNA          HTG          09-DEC-1999
DEFINITION   Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered
            pieces.
ACCESSION   AC018207
VERSION     AC018207.1 GI:6552984
KEYWORDS    HTG; HTGS_PHASE2.
SOURCE      fruit fly.
ORGANISM    Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            Prerygota; Neoptera; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

```

```

REFERENCE   1 (bases 1 to 44887)
AUTHORS     Adams,M. and Venter,J.C.
TITLE       Direct Submission
JOURNAL     Submitted (09-DEC-1999) Celera Genomics, 45 West Gude Drive,
            Rockville, MD, USA
COMMENT     This sequence was identified as CDM:10213855 by the submitter.
            For more information on this record e-mail to fly@celera.com.
            * NOTE: This is a 'working draft' sequence.
            * This sequence will be replaced
            * by the finished sequence as soon as it is available and
            * the accession number will be preserved.
FEATURES
Source      1..44887
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            /db_xref="taxon:7227"
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ORIGIN
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Quality: 39.00 Length: 9
Ratio: 4.875 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889
alignment_block:
US-08-653-294-13 x AC018207/rev ..
Align seg 1/1 to reverse of: AC018207 from: 1 to: 44887
1 TyrArgLeuAlaIleArgLeuAspGlu 9
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17401 TATAGACTGCTATCAGTGCATGAG 17375
seq_name: gb_in2:AC005268
seq_documentation_block:
LOCUS       AC005268      89791 bp      DNA          INV          07-JUL-1998
DEFINITION   Drosophila melanogaster DNA sequence (Pls DS03550 (D225) and
            DS02397 (D206)), complete sequence.
ACCESSION   AC005268 AC004330 AC004314 AC003599
VERSION     AC005268.1 GI:3293207
KEYWORDS    HTG.
SOURCE      Drosophila melanogaster (Subclones in sac from Pl clones DS03550
            (D225) and DS02397 (D206)) DNA.
ORGANISM    Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            Prerygota; Neoptera; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE   1 (bases 1 to 89791)
AUTHORS     Celniker,S.E., George,R.A., Galle,R.F., Hoskins,R.A.,
            Svirskas,R.R., Harris,N.L., Agbayani,A., Arcaina,T.T., Baxter,E.,
            Blazej,R.G., Chavez,C., Chew,M., Doyle,C.M., Farfan,D.E.,
            Flanagan,J., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
            Kim,S.H., Lee,B., Lomotan,M.A., Mak,J.J., Mazda,P., Mok,M.S.,
            Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
            Pfeiffer,B., Punch,E., Snir,E., Twomey,B., Wan,K.H., Whitelaw,K.R.,
            Yee,A., Zhang,R., Zieran,L.L. and Kimmel,B.
            Sequencing of Drosophila chromosome, region 57B6-57C1
            Unpublished (1997)
2 (bases 1 to 89791)
AUTHORS     Celniker,S.E., George,R.A., Galle,R.F., Hoskins,R.A.,
            Svirskas,R.R., Harris,N.L., Agbayani,A., Arcaina,T.T., Baxter,E.,
            Blazej,R.G., Chavez,C., Chew,M., Doyle,C.M., Farfan,D.E.,
            Flanagan,J., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
            Kim,S.H., Lee,B., Lomotan,M.A., Mak,J.J., Mazda,P., Mok,M.S.,
            Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
            Pfeiffer,B., Punch,E., Snir,E., Twomey,B., Wan,K.H., Whitelaw,K.R.,
            Yee,A., Zhang,R., Zieran,L.L. and Kimmel,B.
            Direct Submission
            Submitted (07-JUL-1998) Berkeley Drosophila Genome Project, MS
            64-121, Lawrence Berkeley National Laboratory, One Cyclotron Road,
            Berkeley, CA 94720, US
            Sequence submitted by:

```

Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive web site (<http://fruitfly.berkeley.edu/sequence/>) or send
email to drosophila@hgsc.lbl.gov.
Library locations: 137-37, 136-25.

FEATURES

source
1..89791
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/chromosome="2R"
/map="57B6-57C1"
/clone="pis DS03550 (D225) and DS03397 (D206)"
/note="DS03550 (D225) extends from bp 1 to bp 16,924 and
DS03397 (D206) extends from bp 13,480 to bp 89,791."

BASE COUNT 24290 a 20016 c 20807 g 24678 t
ORIGIN

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Quality: 39.00 Length: 9
Ratio: 4.875 Gaps: 0
Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:

US-08-653-294-13 x AC005268 ..

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24440 TATAGACTGGTATCAGTGCATGAG 24466

seq_name: gb_bal:AMPEPSYNT

seq_documentation_block:
LOCUS AMPEPSYNT 3975 bp DNA BCT 14-OCT-1997
DEFINITION Amycolatopsis mediterranei aps gene, partial.

ACCESSION X97860

VERSION X97860.1 GI:2576250

KEYWORDS APS gene; peptide synthetase.

SOURCE Amycolatopsis mediterranei.

ORGANISM Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;

Actinomycetales; Pseudonocardiaceae; Pseudonocardiaceae;

Amycolatopsis.

REFERENCE 1 (bases 1 to 3975)

AUTHORS Pelzer,S., Reichert,W., Huppert,M., Heckmann,D. and Wohllleben,W.

TITLE Cloning and analysis of a peptide synthetase gene of the balhimycin

producer Amycolatopsis mediterranei DSM5908 and development of a

gene disruption/replacement system

J. Biotechnol. 56 (2), 115-128 (1997)

79749857

REFERENCE 2 (bases 1 to 3975)

AUTHORS Pelzer,S.

TITLE Direct Submission

Submitted (15-MAY-1996) S. Pelzer, Universitaet Tuebingen,

Lehrstuhl Mikrobiologie-Biotechnologie, Auf der Morgenstelle 28, D-

72076 Tuebingen, FRG

Revised by author 14-OCT-97

On Oct 30, 1997 this sequence version replaced gi:1483198.

FEATURES

source

1..3975
/organism="Amycolatopsis mediterranei"

/strain="DSM 5908"

/db_xref="taxon:33910"

gene

1..3975
/gene="aps"

CDS

<1..>3975
/gene="aps"

/codon_start=3

/transl_table=11
/product="peptide-synthetase"
/protein_id="CAA66454.1"
/db_xref="GI:2576251"
/db_xref="SPTREMBL:Q44103"
/translation="IPLDRGEPELTAGDITGLPLRLATLVADTTLVLVHHVVDGW
SAGVEERELAEFYTAAREGPELPELTQYAGYAAEAHVDDQLAYWREQLQAGPR
LAVPTDREAPAQDFAGTREFAPAGLAARIGELAEEDATPFVQAAFAALNRY
TGADLVVGTPTVTRDRPELADLIGYFVNLPRLKIDRAASFRLDVEHVRDTAFDAY
ACLDVDFDVVDALALETPRHPALVQVFGAHEADPAFLREGPLTARRVHHNGTSK
EDFTWSITDDSELGEVEYRSLDEAAVDMTHWRALLTAVLSEPSPLWKIDLEP
VMPVATASQPCRLHSEFSDVDFPFPVAVTFGGASVYIAELDRANRLAHLAEAG
VRPGRVGLLDRLDAIVAILAVLAKAGAAVVPDPAAPDDRAAEVFGDTGRLVYVTD
QCTDGPVDFDLARDVSAYSADRPVPRVPGDYLAIITSGTGRPKGVVAHEHAGR
LLASGHAHGFETEDVWTLFHSAGDWTWELWGLPHHGGRLVLPYLVSRSEAFSTA
LLADGVTMLCQTPSALQLELTATTPRALPALQVLMGGEALDPAVVRWFHASS
APLCNLYGITEVTHVTHDYPGPGAGFERSLIGTLPPLSAHLVLEWLRLPCAGVGE
LYIGGALAHGTWGRAGTAQORFLPDPFSPVPGARLYRTGVARRLAGGLGYVGRCD
SQVKIRGFRIELGEIEHALGAPVACAVTVHDDRLAAVYTGDPDAELRAHLAKSL
PEHMPATVTVLDRLPVTNKGKLDRAALPAPAPRAANAYTAPSTGERLLTLEWSDVL
GVPGAGVHDTNFHLGGDSITRAVHLAKLRDRGWTFTLPDLCAPTAAALPLKPCAG
ETPASRPFAGLSEKDLAKLPQDVVDAYPMAAMQLGMYHMLSGDAGGVHNVSYRVA
GRLDERALRAAVAGAIARHPVLTFTFDVIGYQPMQLVHAEPAPVETADLGLSEA
QRDAVEFDGLCAVRFDLPTPLPFRVQAQLADDDVQLITAEHSHLIDGWSFTSILLT
EILERHADFPDAPPAPPPASTFDFVAEQAAVSAESAERFMRDRLTGANGALWSSGTG
SAATAAEIPTLRLVLPDPAQALAAIAAAGVPAKAVGLAAHRAALALATIGDRVTG
LSVNGRLERSGTAYGLFNTVPLVVDCTERDLVRSVHDEVALLPHRRVFPFARLAR
LMAGPRLCAACFAFLRFHALGRLAGSATSIVDDRIGCEPDMRYEPTNFALVALVQDPA
SGRI"

BASE COUNT 504 a 1504 c 1425 g 542 t

ORIGIN

alignment_scores:

Quality: 38.00 Length: 10
Ratio: 4.222 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-13 x AMPEPSYNT ..

Align seg 1/1 to: AMPEPSYNT from: 1 to: 3975

1 TyrArgLeuAlaIleArgLeuAspGluArg 10

|||||
2907 TACCGGTGCGCGCGCCCTCGACGACGC 2936

seq_name: gb_pr2:HS204E5

seq_documentation_block:

LOCUS HS204E5 129969 bp DNA PRI 22-NOV-1999

DEFINITION Human DNA sequence from PAC 204E5 on chromosome 12. Contains exon
similar to Wilms' Tumour-related protein OM-like P2X-like receptor,
ATP ligand gated ion channel, ESTs, CpG island.

ACCESSION 298941

VERSION 298941.1 GI:2370071

KEYWORDS 12; ATP ligand gated ion channel; CpG island; P2X-like receptor;

OM-like; Tumour-related protein.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 129969)

AUTHORS Kershaw,J.

TITLE Direct Submission

JOURNAL Submitted (29-AUG-1997) Chromosome 12 Project Group

(<http://www.sanger.ac.uk/HGP/Chr12/>) Sanger Centre, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquires:

humquery@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk

IMPORTANT: This sequence is the entire insert of clone 204E5.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together


```

/note="MIR repeat: matches 144. .43 of consensus"
20690. .20979
/note="AluX repeat: matches 302. .10 of consensus"
21716. .22001
/note="AluSx repeat: matches 1. .303 of consensus"
22137. .22350
/note="AluJb repeat: matches 85. .302 of consensus;
incomplete repeat"
23113. .23418
/note="AluJb repeat: matches 302. .1 of consensus"
23420. .23605
/note="MIR repeat: matches 207. .14 of consensus"
23635. .23704
/note="MIR2 repeat: matches 128. .59 of consensus"
23773. .23838
/note="MIR2 repeat: matches 81. .146 of consensus"
24258. .24553
/note="AluYb repeat: matches 1. .301 of consensus"
24560. .24867
/note="AluYb8 repeat: matches 1. .308 of consensus"
24896. .25041
/note="FLAM.A repeat: matches 133. .1 of consensus"
25407. .25708
/note="Alu repeat: matches 303. .2 of consensus"
26138
/note="this base could be T"
26292. .26447
/note="L1 repeat: matches 1943. .1788 of consensus"
26469. .26770
/note="Alu repeat: matches 302. .1 of consensus"
26847. .27145

```

```

alignment_scores:
  Quality: 38.00      Length: 10
  Ratio: 3.800       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 60.000

```

```
alignment_block:
US-08-653-294-13 x HS204E5

```

```
Align seg 1/1 to: HS204E5 from: 1 to: 129969
```

```

1 TyArgLeuAlaIleArgLeuAspGluArg 10
|||||:|||||:|||||:|||||:|||||
71948 TACAGGTAGCAGTCAGATGGACACAGG 71977

```

```
seq_name: gb_pr4:AC009239
```

```

seq_documentation_block:
LOCUS AC009239 136371 bp DNA PRI 22-OCT-1999
DEFINITION Homo sapiens clone NH0470K20, complete sequence.
ACCESSION AC009239
VERSION AC009239.3 GI:6094637
KEYWORDS HTG.
SOURCE human.

```

```

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

```

```

REFERENCE
AUTHORS Waterston,R.H.
TITLE The sequence of Homo sapiens clone
JOURNAL Unpublished
REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (06-AUG-1999) Genome Sequencing Center, Washington

```

```

REFERENCE
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (22-OCT-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
3 (bases 1 to 136371)

```

```

MO 63108, USA
COMMENT On Oct 22, 1999 this sequence version replaced gi:5732147.
FEATURES
Location/Qualifiers
Source
1..136371
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="NH0470K20"

```

```

BASE COUNT 39683 a 25507 c 25721 g 45460 t
ORIGIN

```

```

alignment_scores:
  Quality: 38.00      Length: 9
  Ratio: 4.222       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

```

```
alignment_block:
US-08-653-294-13 x AC009239

```

```
Align seg 1/1 to: AC009239 from: 1 to: 136371
```

```

1 TyArgLeuAlaIleArgLeuAspGlu 9
|||||:|||||:|||||:|||||:|||||
124410 TATAAGTTGCAATTAGATGGATGAA 124436

```

```
seq_name: gb_htg4:AC011830
```

```

seq_documentation_block:
LOCUS AC011830 166832 bp DNA HTG 20-OCT-1999
DEFINITION Homo sapiens chromosome 8 clone 94_A_14 map 8, *** SEQUENCING IN
PROGRESS ***, 17 unordered pieces.
ACCESSION AC011830
VERSION AC011830.2 GI:6087976
KEYWORDS HTG; HTGS_PHASE1.
SOURCE human.

```

```

ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

```

```

REFERENCE
AUTHORS Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE 1 (bases 1 to 166832)
JOURNAL Homo sapiens chromosome 8, clone 94_A_14
REFERENCE
AUTHORS Unpublished

```

```

2 (bases 1 to 166832)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Castle,A., Colangelo,M., Collins,S., Collins,A.,
Cooke,P., Dearellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M.,
Ferrelira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D.,
Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Lehoczky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

```

```

Direct Submission
Submitted (15-OCT-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Oct 20, 1999 this sequence version replaced gi:6041961.
All repeats were identified using RepeatMasker: Smit, A.F.A. &
Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html.

```

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 17 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved. 845: contig of 845 bp in length

```


* 846 2883: contig of 2038 bp in length
* gap of unknown length
* 2884 5685: contig of 2802 bp in length
* gap of unknown length
* 5686 9996: contig of 4311 bp in length
* gap of unknown length
* 9997 14822: contig of 4826 bp in length
* gap of unknown length
* 14823 19370: contig of 4548 bp in length
* gap of unknown length
* 19371 24594: contig of 5224 bp in length
* gap of unknown length
* 24595 30146: contig of 5552 bp in length
* gap of unknown length
* 30147 35542: contig of 5396 bp in length
* gap of unknown length
* 35543 44699: contig of 9157 bp in length
* gap of unknown length
* 44700 55420: contig of 10721 bp in length
* gap of unknown length
* 55421 67431: contig of 12011 bp in length
* gap of unknown length
* 67432 84061: contig of 16630 bp in length
* gap of unknown length
* 84062 100053: contig of 15992 bp in length
* gap of unknown length
* 100054 123949: contig of 23896 bp in length
* gap of unknown length
* 123950 161245: contig of 37296 bp in length
* gap of unknown length
* 161246 166832: contig of 5587 bp in length.
* Location/Qualifiers
* 1..166832
* /organism="Homo sapiens"
* /db_xref="taxon:9606"
* /map="8"
* /clone="94_A_14"
* /clone_lib="RP11 Human Male BAC"
BASE COUNT 45111 a 39419 c 39077 g 42526 t 699 others
ORIGIN

alignment_scores:
Quality: 38.00 Length: 9
Ratio: 4.222 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-08-653-294-13 x AC011830 ..

Align seg 1/1 to: AC011830 from: 1 to: 166832

1 TyrArgLeuAlaIleArgLeuAsglu 9

91099 TACAGGCTGAGCATCCGAGTGATGAG 91125

seq_name: gb_htg3:AC010086

seq_documentation_block:

LOCUS AC010086 187447 bp DNA HTG 04-OCT-1999
DEFINITION Homo sapiens clone NH0209111, *** SEQUENCING IN PROGRESS ***, 3
unordered pieces.

ACCESSION AC010086

VERSION AC010086.2 GI:6007895

KEYWORDS HTG; HTGS_PHASE1.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

Waterston,R.H.

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 187447)
Waterston,R.H.
Direct Submission

Submitted (11-SEP-1999) Genome Sequencing Center, Washington
University School of Medicine, 444 Forest Park Parkway, St. Louis,
MO 63108, USA

On Oct 4, 1999 this sequence version replaced gi:5870314.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 26045: contig of 26045 bp in length
* 26046 26063: gap of unknown length
* 26064 72624: contig of 46561 bp in length
* 72625 72642: gap of unknown length
* 72643 187447: contig of 114805 bp in length.

FEATURES
source
1..187447
Location/Qualifiers

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="NH0209111"

BASE COUNT 56242 a 35522 c 36150 g 59497 t 36 others
ORIGIN

alignment_scores:

Quality: 38.00 Length: 9
Ratio: 4.222 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 77.778

alignment_block:

US-08-653-294-13 x AC010086/rev ..

Align seg 1/1 to reverse of: AC010086 from: 1 to: 187447

1 TyrArgLeuAlaIleArgLeuAsglu 9

69009 TATAAGTTCGATTCGATGATGATGAA 68983

seq_name: gb_htg4:AC008591

seq_documentation_block:

LOCUS AC008591 211769 bp DNA HTG 31-OCT-1999
DEFINITION Homo sapiens chromosome 5 clone CIT-HSPC_575N7, *** SEQUENCING IN
PROGRESS ***, 60 unordered pieces.

ACCESSION AC008591

VERSION AC008591.2 GI:6165161

KEYWORDS HTG; HTGS_PHASE1.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

AUTHORS

DOE Joint Genome Institute.

TITLE Sequencing of Human Chromosome 5

JOURNAL Unpublished

REFERENCE 2 (bases 1 to 211769)

AUTHORS DOE Joint Genome Institute.

TITLE Direct Submission

JOURNAL Unpublished

COMMENT

On Oct 31, 1999 this sequence version replaced gi:5686476.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 60 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 518: contig of 518 bp in length
* gap of unknown length
* 519 1379: contig of 861 bp in length
* gap of unknown length
* 1380 1992: contig of 613 bp in length
* gap of unknown length
* 1993 3019: contig of 1027 bp in length
* gap of unknown length
* 3020 4116: contig of 1097 bp in length
* gap of unknown length
* 4117 4468: contig of 352 bp in length
* gap of unknown length
* 4469 4869: contig of 401 bp in length
* gap of unknown length
* 4870 6986: contig of 2117 bp in length
* gap of unknown length
* 6987 8350: contig of 1364 bp in length
* gap of unknown length
* 8351 9169: contig of 819 bp in length
* gap of unknown length
* 9170 10576: contig of 1407 bp in length
* gap of unknown length
* 10577 10743: contig of 167 bp in length
* gap of unknown length
* 10744 11269: contig of 526 bp in length
* gap of unknown length
* 11270 11921: contig of 652 bp in length
* gap of unknown length
* 11922 12533: contig of 612 bp in length
* gap of unknown length
* 12534 12829: contig of 296 bp in length
* gap of unknown length
* 12830 13906: contig of 1077 bp in length
* gap of unknown length
* 13907 14105: contig of 199 bp in length
* gap of unknown length
* 14106 14914: contig of 809 bp in length
* gap of unknown length
* 14915 16489: contig of 1375 bp in length
* gap of unknown length
* 16490 16732: contig of 243 bp in length
* gap of unknown length
* 16733 16933: contig of 201 bp in length
* gap of unknown length
* 16934 17739: contig of 806 bp in length
* gap of unknown length
* 17740 17875: contig of 136 bp in length
* gap of unknown length
* 17876 19318: contig of 1443 bp in length
* gap of unknown length
* 19319 21247: contig of 1929 bp in length
* gap of unknown length
* 21248 22282: contig of 1035 bp in length
* gap of unknown length
* 22283 23232: contig of 950 bp in length
* gap of unknown length
* 23233 24221: contig of 989 bp in length
* gap of unknown length
* 24222 25716: contig of 1495 bp in length
* gap of unknown length
* 25717 27600: contig of 1884 bp in length
* gap of unknown length
* 27601 28647: contig of 1047 bp in length
* gap of unknown length
* 28648 30401: contig of 1754 bp in length
* gap of unknown length
* 30402 31662: contig of 1261 bp in length
* gap of unknown length
* 31663 34613: contig of 2951 bp in length
* gap of unknown length

* 34614 35537: contig of 924 bp in length
* gap of unknown length
* 35538 37384: contig of 1847 bp in length
* gap of unknown length
* 37385 38975: contig of 1591 bp in length
* gap of unknown length
* 38976 42433: contig of 3458 bp in length
* gap of unknown length
* 42434 46186: contig of 3753 bp in length
* gap of unknown length
* 46187 48955: contig of 2769 bp in length
* gap of unknown length
* 48956 52157: contig of 3202 bp in length
* gap of unknown length
* 52158 57534: contig of 5377 bp in length
* gap of unknown length
* 57535 60590: contig of 3056 bp in length
* gap of unknown length
* 60591 66089: contig of 5499 bp in length
* gap of unknown length
* 66090 70445: contig of 4356 bp in length
* gap of unknown length
* 70446 77265: contig of 8820 bp in length
* gap of unknown length
* 77266 82716: contig of 5451 bp in length
* gap of unknown length
* 82717 87967: contig of 5251 bp in length
* gap of unknown length
* 87968 95151: contig of 7184 bp in length
* gap of unknown length
* 95152 102796: contig of 7645 bp in length
* gap of unknown length
* 102797 111023: contig of 8227 bp in length
* gap of unknown length
* 111024 119350: contig of 8327 bp in length
* gap of unknown length
* 119351 129014: contig of 9664 bp in length
* gap of unknown length
* 129015 140858: contig of 11844 bp in length
* gap of unknown length
* 140859 154309: contig of 13451 bp in length
* gap of unknown length
* 154310 164838: contig of 10529 bp in length
* gap of unknown length
* 164839 177853: contig of 13015 bp in length
* gap of unknown length
* 177854 192072: contig of 14219 bp in length
* gap of unknown length
* 192073 211769: contig of 19697 bp in length.
* Location/Qualifiers
1. 211769
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CIT-HSPC-575N7"

BASE COUNT 64860 a 39430 c 40173 g 66628 t 678 others
ORIGIN

alignment_scores:
Quality: 38.00 Length: 10
Ratio: 4.222 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-13 x AC008591/rev ..

Align seg 1/1 to reverse of: AC008591 from: 1 to: 211769

1 TTTATGLeuAlaileArgLeuAspGluArg 10

||||||| :|||:|||||:|||||

30417 TACCGCTATAGTTCGGCTCGACTCTAGA 30388

```

seq_name: gb_in1:AF026152
seq_documentation_block:
LOCUS AF026152 486 bp DNA INV 26-OCT-1997
DEFINITION Caenorhabditis elegans chitinase (CHT1) gene, partial cds.
ACCESSION AF026152
VERSION AF026152.1 GI:2564714
KEYWORDS
SOURCE
ORGANISM
Caenorhabditis elegans.
Caenorhabditis elegans
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE
AUTHORS de la Vega, H., Specht, C.A., Liu, Y. and Robbins, P.W.
TITLE Chitinases are a multi-gene family in Aedes, Anopheles, and
Drosophila
JOURNAL Insect Mol. Biol. (1997) In press
AUTHORS
TITLE
JOURNAL
AUTHORS
TITLE Direct Submission
JOURNAL
SUBMITTED (19-SEP-1997) Biology, Massachusetts Institute of
Technology, 40 Ames St. E17-235, Cambridge, MA 02139, USA
FEATURES
Location/Qualifiers
source
1..486
/organism="Caenorhabditis elegans"
/db_xref="taxon:6239"
join(<1..14,150..>486)
/mrna
/gene="CHT1"
/product="Chitinase"
<1..>486
/gene="CHT1"
CDS
join(<1..14,150..>486)
/gene="CHT1"
/codon_start=1
/product="Chitinase"
/protein_id="AAB81847.1"
/db_xref="GI:2564715"
/translation="WAQYRGRAKFVPEYDPGICLTHLFAFGWMNADYTVRAYDPAD
LPNDWAGEGMYRRYKLVTDLTQLLSFGCSFGTALFGMAASSASRVFIDSAT
TSVRTGFGDGLDW"
BASE COUNT 130 a 90 c 120 g 146 t
ORIGIN

alignment_scores:
Quality: 37.00 Length: 10
Ratio: 4.11 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-13 x AF026152 ..
Align seg 1/1 to: AF026152 from: 1 to: 486
1 TyrArgLeuAlaIleArgLeuAspGluArg 10
|||||:|||||:|||||:|||||
204 TATTCCTTTGCAATCGATGATGAACGC 233

seq_name: gb_ba1:HSOMXXU
seq_documentation_block:
LOCUS HSOMXXU 1323 bp DNA BCT 10-APR-1997
DEFINITION Hyphomicrobium sp. DNA, Mox mutant mxu-1::Tn5-132.
ACCESSION Y09104
VERSION Y09104.1 GI:1869806
KEYWORDS methanol oxidation system; Mox.
SOURCE Hyphomicrobium sp.
ORGANISM
Bacteria; Proteobacteria; alpha subdivision; Hyphomicrobium.
REFERENCE
AUTHORS Gliesche, C.G., Menzel, M. and Fesefeldt, M.
TITLE A rapid method for creating species-specific gene probes for
methylophilic bacteria

```

```

J. Microbiol. Methods 28, 25-34 (1997)
REFERENCE
2 (bases 1 to 1323)
AUTHORS Gliesche, C.G.
TITLE Direct Submission
JOURNAL
SUBMITTED (24-OCT-1996) C.G. Gliesche, CAU-Kiel, Institut fuer
Allgemeine Mikrobiologie, Am Botanischen Garten 1-9,
(Biologisches Zentrum), D-24118 Kiel, FRG
FEATURES
Location/Qualifiers
source
1..1323
/organism="Hyphomicrobium sp."
/strain="B 69"
/db_xref="taxon:82"
<1..>1323
/mrna
/note="mxu-1::Tn5-132; fragment of methanol oxidation
system (Mox)"
BASE COUNT 221 a 402 c 429 g 271 t
ORIGIN

alignment_scores:
Quality: 37.00 Length: 8
Ratio: 4.625 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 87.500

alignment_block:
US-08-653-294-13 x HSOMXXU ..
Align seg 1/1 to: HSOMXXU from: 1 to: 1323
1 TyrArgLeuAlaIleArgLeuAsp 8
|||||:|||||:|||||:|||||
967 TATCGCTGGCGCTTCGTCGAT 990

seq_name: gb_in1:CELC04F6
seq_documentation_block:
LOCUS CELC04F6 25083 bp DNA INV 19-DEC-1995
DEFINITION Caenorhabditis elegans cosmid C04F6.
ACCESSION U42835
VERSION U42835.1 GI:1125760
KEYWORDS
SOURCE
Caenorhabditis elegans strain=Bristol N2.
Caenorhabditis elegans
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
REFERENCE
1 (bases 1 to 25083)
AUTHORS Wilson, R., Ainscough, R., Anderson, K., Baynes, C., Berks, M.,
Bonfield, J., Burton, J., Connell, M., Copsey, T., Cooper, J.,
Coulson, A., Craxton, M., Dear, S., Du, Z., Durbin, R., Favello, A.,
Fulton, L., Gardner, A., Green, P., Hawkins, T., Hillier, L., Jier, M.,
Johnston, L., Jones, M., Kersey, J., Kirsten, J., Laister, N.,
Latreille, P., Lightning, J., Lloyd, C., McMurray, A., Mortimore, B.,
O'Callaghan, M., Parsons, J., Percy, C., Riffen, L., Roopra, A.,
Saunders, D., Showkhen, R., Smalton, N., Smith, A., Sounhammer, E.,
Staden, R., Sulston, J., Thierry-Mieg, J., Thomas, K., Vaudin, M.,
Vaughan, K., Waterston, R., Watson, A., Weinstock, L.,
Walkinson-Sproat, J. and Wohldman, P.
2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans
elegans
Nature 368 (6466), 32-38 (1994)
JOURNAL
MEDLINE 94150718
REFERENCE
2 (bases 1 to 25083)
AUTHORS Nhan, M.
TITLE The sequence of C. elegans cosmid C04F6
JOURNAL Unpublished (1995)
REFERENCE
3 (bases 1 to 25083)
AUTHORS Waterston, R.
TITLE Direct Submission
JOURNAL Submitted (10-DEC-1995) Robert Waterston
COMMENT Submitted by:
Genome Sequencing Center
Department of Genetics, Washington University,
St. Louis, MO 63110, USA, and

```

Sanger Centre, Hinxton Hall
 Cambridge CB10 1RQ, England
 e-mail: rwenemato@wustl.edu and jesus@anger.ac.uk
 NEIGHBORING COSMID INFORMATION:

The 5' cosmid is ZK563, 200 bp overlap; 3' cosmid is ZK813, 200 bp overlap. Actual start of this cosmid is at base position 197 of CELC04F6; actual end is at 20314 of CELZK813

NOTES:

Coding sequences below are predicted from computer analysis, using the program Genefinder (P. Green and L. Hillier, ms in preparation).

FEATURES

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About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 CompuGen Ltd.

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	N_Geneseq_36:T22674	-	31.00	69.14	2.9e+04	50341	Bacillus subtilis srfA operon
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	N_Geneseq_36:T51411	-	31.00	68.80	3.0e+04	52297	Mycobacteriophage L5 genome s

N_Geneseq_36:047357 - 31.00 68.80 3.0e+04 52298 ! L5 mycobacteriophage DNA.
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AC T80415;
DT 02-MAR-1998 (first entry)
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KW tyactone synthase gene cluster; tylg gene; multifunctional protein;
KW platenolide synthase gene cluster; platenolide production; smtg gene;
KW polyketide; tyactone synthesis; antibiotic; tylosin; hybrid gene; ss.
OS Streptomyces ambofaciens.
OS Streptomyces fradiae.
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PD 27-AUG-1997.
PF 13-FEB-1997; 301056.
PR 22-FEB-1996; US-012078.
PA (ELIL) LILLY & CO ELI.
PI Dehoff BS, Kuhstoss SA, Rostock PR, Sutton KL;
DR WPI; 97-418046/39.
DR P-PSDB; W22611.
PT DNA encoding Streptomyces fradiae tyactone synthase domain - for
PT production of tylosin-related polyketide compounds
PS Claim 22; Pages 178-197; 220pp; English.
CC This sequence represents a hybrid gene of the invention. This sequence
CC was created by replacing a EcoRI-ApaI fragment of smtg ORF1 with a
CC EcoRI-StuI fragment from tylg ORF1. The position of the nucleotides from
CC each of the two genes is not given in the specification. The smtg gene
CC (see T80414) was isolated from Streptomyces ambofaciens, and encodes the
CC multi-functional proteins which direct the synthesis of the polyketide
CC platenolide. Platenolide is the basic building block of the macrolide
CC antibiotic spiramycin. The tylg gene (see T80413) is the tyactone
CC synthase gene cluster of the invention. The tylg sequence was isolated
CC from Streptomyces fradiae, and encodes multifunctional proteins which
CC direct the synthesis of the polyketide tyactone. Tyactone is the basic
CC building block of the antibiotic tylosin. The hybrid sequence can be used
CC to transform S. ambofaciens lacking the smtg ORF1 sequence, or S. fradiae
CC lacking the tylg ORF1 sequence, so that they can produce polyketides. The
CC DNA sequence can be modified so as to alter the type of carboxylic acids
CC incorporated, the number of carboxylic acids incorporated and/or the
CC post-condensation reactions performed, thereby resulting in novel
CC tylosin-related polyketides.
SQ Sequence 13987 BP; 1556 A; 4401 C; 5727 G; 2303 T;

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AC T78508;

DT 26-FEB-1998 (first entry)
 DE Platenolide synthase gene cluster.
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 KW multi-functional protein; macrolide antibiotic; spiramycin; ss.
 OS Streptomyces ambofaciens.
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 FT /note= "ORF4 encodes protein shown in W23719"
 CDS 36155..41830
 FT /tag= e
 FT /note= "ORF5 encodes protein shown in W23720"
 PN EP-791655-A2.
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 PF 19-FEB-1997; 301066.
 PR 22-FEB-1996; US-012050.
 PA (ELIL) LILLY & CO ELI.
 PI Burgett SG, Kuhstoss SA, Rao RN, Richardson MA;
 PI Rosteck PR;
 DR WPI: 97-418047/39.
 DR P-PSDB; W23716-W23720.
 PT DNA encoding Streptomyces ambofaciens platenolide synthase domain -
 PT for production of spiramycin-related polyketide antibiotics
 PS Claim 9; Pages 8-33; 81pp; English.
 CC This sequence represents the platenolide synthase gene cluster of the
 CC invention. This sequence is referred to as the smg gene, and was
 CC isolated from Streptomyces ambofaciens. This sequence encodes the
 CC multi-functional proteins which direct the synthesis of the polyketide
 CC platenolide. Platenolide is the basic building block of the macrolide
 CC antibiotic spiramycin. The DNA can be used to produce compounds
 CC exhibiting antibiotic activity based on the platenolide structure,
 CC including specifically the macrolide antibiotic spiramycin and spiramycin
 CC analogues and derivatives. Modifications of the platenolide synthase DNA
 CC sequence can be made so as to change the number and type of carboxylic
 CC acids incorporated into the growing polyketide chain and to change the
 CC kind of post-condensation processing that is conducted.
 SQ Sequence 44377 BP; 4965 A; 15552 C; 17381 G; 6479 T;

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 US-08-653-294-13 x T78508/rev ..
 Align seg 1/1 to reverse of: T78508 from: 1 to: 44377

1 TyrArgLeuAlaIleArgLeuAspGlu 9
 |||||
 6667 TACCGCTCGCCAGCGCTGGACGAG 6641

seq_name: N_Geneseq_36:T80414

seq_documentation_block:

ID T80414 standard; DNA; 44377 BP.
 AC T80414;
 DT 27-FEB-1998 (first entry)
 DE Platenolide synthase gene cluster.
 KW Tylactone synthase gene cluster; tylG gene; multifunctional protein;
 KW platenolide synthase gene cluster; platenolide production; smg gene;

KW polyketide; tylactone synthesis; antibiotic; tylosin; ss.
 OS Streptomyces ambofaciens.
 FH Key Location/Qualifiers
 FT 350..14002
 FT /tag= a
 FT /transl_except= (pos:350..352, aa:Met)
 FT /note= "ORF1 encodes protein shown in W22606"
 CDS 14046..20036
 FT /tag= b
 FT /note= "ORF2 encodes protein shown in W22607"
 CDS 20110..31284
 FT /tag= c
 FT /transl_except= (pos:20111..20113, aa:Met)
 FT /note= "ORF3 encodes protein shown in W22608"
 CDS 31329..36071
 FT /tag= d
 FT /note= "ORF4 encodes protein shown in W22609"
 CDS 36155..41830
 FT /tag= e
 FT /note= "ORF5 encodes protein shown in W22610"
 PN EP-791655-A2.
 PD 27-AUG-1997.
 PF 19-FEB-1997; 301056.
 PR 22-FEB-1996; US-012078.
 PA (ELIL) LILLY & CO ELI.
 PI Dehoff BS, Kuhstoss SA, Rosteck PR, Sutton KL;
 DR WPI: 97-418046/39.
 DR P-PSDB; W22606-W22610.
 PT DNA encoding Streptomyces fradiae tylactone synthase domain - for
 PT production of tylosin-related polyketide compounds
 PS Example 2; Pages 110-134; 220pp; English.
 CC This sequence represents the platenolide synthase gene cluster of the
 CC invention. This sequence is referred to as the smg gene, and was
 CC isolated from Streptomyces ambofaciens. This sequence encodes the
 CC multi-functional proteins which direct the synthesis of the polyketide
 CC platenolide. Platenolide is the basic building block of the macrolide
 CC antibiotic spiramycin. This sequence was used along with the tylG gene
 CC is the tylactone synthase gene cluster of the invention. The tylG gene
 CC sequence was isolated from Streptomyces fradiae, and encodes
 CC multifunctional proteins which direct the synthesis of the polyketide
 CC tylactone. Tylactone is the basic building block of the antibiotic
 CC tylosin. The hybrid sequence can be used to transform S. ambofaciens
 CC lacking the smg ORF1 sequence, or S. fradiae lacking the tylG ORF1
 CC sequence, so that they can produce polyketides. The DNA sequence can be
 CC modified so as to alter the type of carboxylic acids incorporated, the
 CC number of carboxylic acids incorporated and/or the post-condensation
 CC reactions performed, thereby resulting in novel tylosin-related
 CC polyketides.
 SQ Sequence 44377 BP; 4965 A; 15552 C; 17381 G; 6479 T;

alignment_scores:
 Quality: 37.00 Length: 9
 Ratio: 4.625 Gaps: 0
 Percent Similarity: 88.889 Percent Identity: 88.889

alignment_block:

US-08-653-294-13 x T80414/rev ..
 Align seg 1/1 to reverse of: T80414 from: 1 to: 44377

1 TyrArgLeuAlaIleArgLeuAspGlu 9
 |||||
 6667 TACCGCTCGCCAGCGCTGGACGAG 6641

seq_name: N_Geneseq_36:T04154

seq_documentation_block:

ID T04154 standard; DNA; 1791 BP.
 AC T04154;
 DT 26-FEB-1996 (first entry)
 DE BCG DapB gene.

KW Dihydrodipicolinate-reductase; diaminopimelate-dehydrogenase; dapB;
 KW diaminopimelic acid; peptidoglycan; mycobacteria; vaccine;
 KW Mycobacterium tuberculosis; Mycobacterium avium;
 KW Mycobacterium fortuitum; Mycobacterium leprae; Mycobacterium goodii;
 KW Mycobacterium haemophilum; Mycobacterium paratuberculosis; BCG; ss.
 OS Mycobacterium bovis.
 FH Key Location/Qualifiers
 FT rbs 297..303
 FT rbs /*tag= a
 FT cds 312..1127
 FT /*tag= b
 FT /product= DapB
 FT rbs 1136..1140
 FT /*tag= c
 FT cds 1151..1683
 FT /*tag= d
 FT /product= unidentified
 PN W09532226-A1.
 PD 31-AUG-1995.
 PF 27-FEB-1995; U02455.
 PR 28-FEB-1994; US-203190.
 PA (YESH) UNIV YESHIVA EINSTEIN COLLEGE.
 PI Cirillo JD, Jacobs WR.
 DR WPI; 95-311535/40.
 DR P-PDB; R79946.
 PT Gene involved in the synthesis of di-aminopimelic acid in
 PT mycobacteria - and methods for inhibiting its expression to treat
 PT mycobacterial infection
 PS Claim 3: Fig 3: 48pp; English.
 CC A BCG DNA fragment (T04154) that complemented a dapB mutation in
 CC Escherichia coli was isolated. The BCG dapB gene encoded a
 CC bifunctional enzyme (R79946) capable of catalyzing the
 CC dihydrodipicolinate-reductase and diaminopimelate-dehydrogenase
 CC reactions involved in diaminopimelic acid (DAP) biosynthesis. DAP
 CC biosynthetic genes are useful as targets for anti-mycobacterial
 CC agents and for the design of in vivo selection systems. Mycobacteria
 CC having a deleted dapB gene can be used for vaccine prodn.
 SQ Sequence 1791 BP; 254 A; 638 C; 640 G; 259 T;

alignment_scores:
 Quality: 34.00 Length: 9
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 66.667

alignment_block:

US-08-653-294-13 x T04154/rev ..

Align seg 1/1 to reverse of: T04154 from: 1 to: 1791

1 TyrArgLeuAlaIleArgLeuAspGlu 9

.....
 716 TTTCGTCTCGGATCGCTCAGCAT 690

seq_name: N_Geneseq_36:T06770

seq_documentation_block:

ID T06770 standard; DNA; 4602 BP.
 AC T06770;
 DT 15-OCT-1996 (first entry)
 DE Pseudomonas aureofaciens phenazine gene cluster.
 KW Antipathogenic substance; phenazine; antibiotic;
 KW fungicide; pesticide; ss.
 OS Pseudomonas aureofaciens.
 FH Key Location/Qualifiers
 FT cds 230..1597
 FT /*tag= a
 FT /note= "phz1 (ORF1)"
 FT cds 1598..2761
 FT /*tag= b
 FT /note= "phz2 (ORF2)"
 FT cds 2764..3600
 FT /*tag= c

FT /note= "phz3 (ORF3)"
 FT 3597..4265
 FT /*tag= d
 FT /note= "phz4 (ORF4)"
 PN W09533818-A2.
 PD 14-DEC-1995.
 PF 30-MAY-1995; IB0414.
 PR 08-JUN-1994; US-258261.
 PA (CIBA) CIBA GEIGY AG.
 PI Beck JJ, Gaffney TD, Hammer PE, Hill DS, Lam ST;
 PI Ligon J, Ryals JA, Schupp T, Uknes SJ;
 DR WPI; 96-040226/04.
 DR P-PDB; R87533, R87534, R87535, R87536.
 PT New genes for biosynthesis of anti-pathogenic substances - pref.
 PT pyrolnitrin and soraphen, useful for disease control in plants
 PS Disclosure; Page 163-169; 190pp; English.
 CC This is the phenazine gene cluster encoding 4 open reading frames
 CC which each encode 1 polypeptide. The gene cluster may be expressed
 CC recombinantly to produce phenazine, or expressed in a transgenic
 CC plant for disease-resistance.
 SQ Sequence 4602 BP; 925 A; 1516 C; 1341 G; 818 T;

alignment_scores:
 Quality: 34.00 Length: 10
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:
 US-08-653-294-13 x T06770 ..

Align seg 1/1 to: T06770 from: 1 to: 4602

1 TyrArgLeuAlaIleArgLeuAspGluArg 10
 |||||
 994 TATCGACTTGCTGCCACCTCGATCGGCGC 1023

seq_name: N_Geneseq_36:T89957

seq_documentation_block:
 ID T89957 standard; DNA; 4603 BP.
 AC T89957;
 DT 12-MAR-1998 (first entry)
 DE Pseudomonas aureofaciens phenazine gene cluster genomic DNA.
 KW Phenazine; biosynthesis; antibiotic; antipathogenic;
 KW transgenic plant; phytopathogen; resistance; ss.
 OS Pseudomonas aureofaciens.
 FH Key Location/Qualifiers
 FT cds 230..1597
 FT /*tag= a
 FT /product= phz1
 FT 1598..2761
 FT /*tag= b
 FT /product= phz2
 FT /transl_except= (pos: 1796..1798, aa: Xaa)
 FT /note= "Xaa = unknown"
 FT 2764..3600
 FT /*tag= c
 FT /product= phz3
 FT 3597..4265
 FT /*tag= d
 FT /product= phz4
 FT /transl_except= (pos: 3391..3393, aa: Xaa)
 FT /note= "Xaa = unknown"
 PN US5662898-A.
 PD 02-SEP-1997.
 PF 01-JUN-1995; 457342.
 PR 20-AUG-1990; US-570184.
 PR 02-JUL-1992; US-908284.
 PR 31-AUG-1992; US-937648.
 PR 01-JUL-1993; US-087636.
 PR 08-JUN-1994; US-258261.
 PA (CIBA) CIBA GEIGY CORP.

PI Beck JJ, Gaffney TD, Hammer PE, Hill DS, Lam ST;
 PI Ligon JM, Ryals JA, Schupp T, Uknes SJ;
 DR WPI: 97-447901/41.
 DR P-PSDB: W31304; W31305; W31306; W31307.
 PT Protecting plants against pathogens with genetically transformed
 PT biological control agent - which expresses all polypeptide(s)
 PT involved in pyrrolnitrin biosynthetic pathway
 PS Example 18; Column 135-144; 88pp; English.
 CC This genomic DNA sequence encodes a cluster of genes involved in
 CC phenazine biosynthesis. Phenazines are nitrogen-containing
 CC heterocyclic compounds with a common planar aromatic tricyclic
 CC structure. It has been proposed that phenazine antibiotic function
 CC arises from the formation of intercalative complexes with DNA
 CC interfering with DNA metabolism. Transgenic plants containing such
 CC antipathogenic genes should have enhanced resistance to attack by
 CC phytopathogens.
 SQ Sequence 4603 BP; 925 A; 1516 C; 1342 G; 818 T;
 alignment_scores:
 Quality: 34.00 Length: 10
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000
 alignment_block:
 US-08-653-294-13 x T89957 ..
 Align seg 1/1 to: T89957 from: 1 to: 4603
 1 TyrArgLeuAlaIleArgLeuAspGluArg 10
 ||||| ||||| :||| ||||| :|||
 994 TATCGACTTGCCTGCCACCTCGATCGCGC 1023
 seq_name: N_Geneseq_36:V58732
 seq_documentation_block:
 ID V58732 standard; DNA: 4603 BP.
 AC V58732;
 DT 07-DEC-1998 (first entry)
 DE Phenazine gene cluster
 KW Pyrrolnitrin; biosynthetic pathway; pathogen protection; phenazine;
 KW plant antipathogenic substance production; anti-fungal antibiotic;
 KW fungal respiratory electron transport inhibitor; lipoprotein damage; ss.
 OS Pseudomonas aureofaciens.
 PH Key Location/Qualifiers
 FT CDS 230..1597
 FT /*tag= a
 FT /product= phz1
 FT CDS 1598..2761
 FT /*tag= b
 FT /product= phz2
 FT CDS 2764..3600
 FT /*tag= c
 FT /product= phz3
 FT CDS 3597..4265
 FT /*tag= d
 FT /product= phz4
 US5817502-A.
 PD 06-OCT-1998.
 PF 09-OCT-1998; 729214.
 PR 09-OCT-1998; US-729214.
 PR 08-JUN-1994; US-258261.
 PR 30-MAY-1995; WO-IB0414.
 PA (NOVS) NOVARTIS FINANCE CORP.
 PI Hammer PE, Hill DS, Kirner S, Lam ST, Ligon JM,
 PI Van Pee K.
 DR WPI: 98-536391/47.
 DR P-PSDB: W69401, W69402, W69403, W69404.
 PT Genes encoding enzymes of the biosynthetic pathway of pyrrolnitrin -
 PT useful for producing transgenic plants which can produce
 PT pyrrolnitrin as an anti-pathogenic agent
 PS Example 18; Column 147-156; 109pp; English.
 CC This sequence represents the phenazine gene cluster, isolated from

CC Pseudomonas aureofaciens. This sequence was used to isolate the
 CC pyrrolnitrin gene region of the invention, that encodes at least one
 CC enzyme required in the biosynthetic pathway of pyrrolnitrin. The DNA and
 CC host cells transformed with it are useful for the production of
 CC transgenic plants with protection against phytopathogens. The enzymes
 CC are part of a biosynthetic pathway producing plant antipathogenic
 CC substances (APS). The compound ultimately produced by the pathway,
 CC pyrrolnitrin, is a broad range, anti-fungal antibiotic. It inhibits
 CC fungal respiratory electron transport and causes general lipoprotein
 CC damage. The transformed cells can additionally be used in compositions to
 CC be applied to plants to provide resistance, as can purified APS produced
 CC by them. Transgene dependent resistance eliminates the need to spray
 CC crops with chemical based pesticides and antibiotics, which is expensive
 CC and time consuming, and in addition, especially in the case of
 CC antibiotics, their over use leads to resistance. In addition, transgenic
 CC production of these enzymes overcomes problems of applying
 CC micro-organisms which can be slow growing and isolated in their growth
 CC areas.
 SQ Sequence 4603 BP; 925 A; 1516 C; 1342 G; 818 T;
 alignment_scores:
 Quality: 34.00 Length: 10
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000
 alignment_block:
 US-08-653-294-13 x V58732 ..
 Align seg 1/1 to: V58732 from: 1 to: 4603
 1 TyrArgLeuAlaIleArgLeuAspGluArg 10
 ||||| ||||| :||| ||||| :|||
 994 TATCGACTTGCCTGCCACCTCGATCGCGC 1023
 seq_name: N_Geneseq_36:V39844
 seq_documentation_block:
 ID V39844 standard; DNA: 5698 BP.
 AC V39844;
 DT 29-SEP-1998 (first entry)
 DE Pseudomonas fluorescens phenazine gene cluster.
 KW Pseudomonas; genetic engineering; biocontrol; plant; pathogenic;
 KW Rhizoctonia; Pythium; antifungal; pyrrolnitrin; crop protection; ss.
 OS Pseudomonas fluorescens.
 PH Key Location/Qualifiers
 FT CDS 105..1307
 FT /*tag= a
 FT /product= "phzF"
 FT /note= "ORF1"
 FT CDS 1323..1946
 FT /*tag= b
 FT /product= "phzA"
 FT /note= "ORF2"
 FT CDS 1943..3856
 FT /*tag= c
 FT /product= "phzB"
 FT /note= "ORF3"
 FT CDS 3859..4695
 FT /*tag= d
 FT /product= "phzC"
 FT /note= "ORF4"
 FT CDS 4692..5360
 FT /*tag= e
 FT /product= "phzD"
 FT /note= "ORF5"
 WO9824919-A1.
 PD 11-JUN-1998.
 PF 05-DEC-1997; E06815.
 PR 09-SEP-1997; US-058304.
 PR 06-DEC-1996; US-761258.
 PA (NOVS) NOVARTIS AG.
 PI Gaffney TD, Hill DS, Lam ST, Ligon JM, Stafford JM,

PI Torkewitz NR:
 DR WPI; 98-33337/29.
 PT Genetically modified pseudomonas strains - useful to protect crop
 PT plants by controlling or inhibiting plant pathogen growth, e.g.
 PT growth of Rhizoctonia species
 PS Example 9; Page 71-75; 85pp; English.
 CC A genetically engineered biocontrol strain of Pseudomonas has been
 CC developed that can control attacks on crop plants by pathogenic fungi,
 CC e.g. Rhizoctonia and Pythium and aggressively compete with indigenous
 CC bacteria and microflora in the plant rhizosphere. The strains can be
 CC included with agronomically acceptable carriers or chemical fungicides
 CC (e.g. metalaxyl compounds) in biocontrol compositions. The strains or
 CC compositions can be applied to a plant/plant part to protect it from a
 CC plant pathogenic fungus, by controlling or inhibiting fungal growth.
 CC They can also be applied to the environment in which a plant pathogen
 CC fungus will grow (e.g. soil) to similarly control or inhibit pathogen
 CC growth, or to seeds to protect plants developing from the seed from a
 CC plant pathogenic fungus. They are especially effective against
 CC Rhizoctonia and Pythium species which cause damping off in cotton.
 CC Rhizoctonia also infects many other crop species (e.g. beans and wheat),
 CC and no effective chemical fungicides are available. The present sequence
 CC represents the Pseudomonas fluorescens phenazine gene cluster used
 CC in an example from the present invention.
 SQ Sequence 5698 BP; 1166 A; 1855 C; 1677 G; 1000 T;

alignment_scores:
 Quality: 34.00 Length: 10
 Ratio: 3.778 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-13 x V39844 ..

Align seg 1/1 to: V39844 from: 1 to: 5698

1 TyrArgLeuAlaIleArgLeuAspLarg 10

||||| :||| :||| :|||

2089 TATCGACTGCTGCGCAGCTGATCGGCGC 2118

seq_name: N_Geneseq_36:X51895

seq_documentation_block:

ID X51895 standard; DNA; 278 BP.

AC X51895;

DT 22-JUN-1999 (first entry)

DE Human secreted protein 5', EST SEQ ID NO: 109.

KW Human; secreted protein; EST; expressed sequence tag; diagnosis;

KW forensic; gene therapy; chromosome mapping; signal peptide;

KW upstream regulatory sequence; cytokine activity; cell proliferation;

KW differentiation; haematopoiesis regulation; tissue growth regulation;

KW reproductive hormone regulation; chemotactic; chemokinetic; haemostatic;

KW thrombolytic; anti-inflammatory; tumour inhibition; ds.

OS Homo sapiens.

PN WO9906552-A2.

PF 31-JUL-1998; IB1236.

PP 11-FEB-1999.

PR 01-AUG-1997; US-905223.

PA (GEST) GENSET.

PI Duclert A, Dumas Milne Edwards J, Lacroix B;

DR WPI; 99-153782/13.

DR P-PSDB; Y13095.

PT New isolated brain-derived nucleic acids - used to develop products

PT which may have cytokine, immune, regulatory, haematopoiesis

PT regulating, anti-inflammatory or tumour inhibition activity

PS Claim 1; Page 236; 577pp; English.

CC X51787 to X52019 represent 5' expressed sequence tags (ESTs) for human

CC secreted proteins, and encode the proteins given in Y1287 to Y13219,

CC respectively. The proteins given represent the signal peptide and an

CC N-terminal fragment of a secreted protein. The nucleic acid sequences

CC can be used for producing secreted human gene products. They can also

CC be used to develop products for diagnosis and therapy. The proteins

CC obtained may have cytokine activity, cell proliferation/differentiation

CC activity, haematopoiesis regulating activity, tissue growth regulating
 CC activity, reproductive hormone regulating activity, chemotactic/
 CC chemokinetic activity, haemostatic and thrombolytic activity, receptor/
 CC ligand activity, anti-inflammatory activity, tumour inhibition activity
 CC or other activities. The products can be used in forensic, gene therapy
 CC and chromosome mapping procedures. The sequences can also be used for
 CC obtaining corresponding promoter sequences. The nucleic acids encoding
 CC the signal peptide can be used for directing extracellular secretion of
 CC a polypeptide or the insertion of a polypeptide into a membrane, or
 CC importing a polypeptide into a cell.
 SQ Sequence 278 BP; 95 A; 56 C; 58 G; 67 T;

alignment_scores:

Quality: 33.00 Length: 9

Ratio: 4.125 Gaps: 0

Percent Similarity: 88.889 Percent Identity: 66.667

alignment_block:

US-08-653-294-13 x X51895 ..

Align seg 1/1 to: X51895 from: 1 to: 278

1 TyrArgLeuAlaIleArgLeuAspLarg 9

||||| :||| :||| :|||

241 TACAGACTAGCTGCGCATCAGAG 267

seq_name: N_Geneseq_36:V74662

seq_documentation_block:

ID V74662 standard; DNA; 1816 BP.

AC V74662;

DT 16-MAR-1999 (first entry)

DE Staphylococcus aureus contig SEQ ID #351.

KW Computer readable medium; vaccine; S.aureus infection; immunodetection;

KW cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;

KW skin infection; surgical wound infection; scalded skin syndrome;

KW toxic shock syndrome; ds.

OS Staphylococcus aureus.

PH Key Location/Qualifiers

FT misc_feature 241..300

FT /*tag= a

FT /note= "these bases represent a line of missing text in
 the sequence listing in the specification. They
 are included to maintain the nucleotide numbering
 given in the specification for this DNA sequence"

FT

FT

FT

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FT

CC (and their fragments) are useful as primers or probes for isolating
 CC homologues of any of the S.aureus DNA sequences contained on the
 CC computer readable medium.
 SQ Sequence 1816 BP; 539 A; 307 C; 240 G; 667 T;

alignment_scores:
 Quality: 33.00 Length: 9
 Ratio: 4.14 Gaps: 0
 Percent Similarity: 77.78 Percent Identity: 77.78

alignment_block:

US-08-653-294-13 x V74662/rev ..

Align seg 1/1 to reverse of: V74662 from: 1 to: 1816

1 TyrArgLeuAlaIleArgLeuAspGlu 9

1387 TATCGTTTACTAATAACATTAGACGAA 1361

seq_name: N_Geneseq_36.T15599

seq_documentation_block:

ID T15599 standard; DNA; 2880 BP.

AC T15599;

DT 07-APR-1996 (first entry)

DE Laccase-LCC2 gene.

KW Laccase-LCC2: Polyporus pinsitus; Trametes villosa; primer;

KW polymerase chain reaction; PCR; signal peptide; cellulase;

KW Aspergillus oryzae; cDNA probe; Escherichia coli; plasmid pDSV19;

KW lignin; lignosulphonate; polymerisation; Kraft pulp;

KW depolymerisation; oxidation; hair dye; phenol; aniline; vector;

KW cloning; basidiomycetes; ss.

OS Polyporus pinsitus.

OS Trametes villosa.

PH Key Location/Qualifiers

FT cds 364..2492

FT /*tag= a

FT /product= Laccase-LCC2

FT /note= "PC-1.10.3.2"

FT signal_peptide 364..423

FT /*tag= b

FT exon 364..543

FT /*tag= c

FT intron 544..592

FT /*tag= d

FT exon 593..661

FT /*tag= e

FT intron 662..715

FT /*tag= f

FT exon 716..835

FT /*tag= g

FT intron 836..899

FT /*tag= h

FT exon 900..1013

FT /*tag= i

FT intron 1014..1066

FT /*tag= j

FT exon 1067..1132

FT /*tag= k

FT intron 1133..1187

FT /*tag= l

FT exon 1188..1283

FT /*tag= m

FT intron 1284..1343

FT /*tag= n

FT exon 1344..1498

FT /*tag= o

FT intron 1499..1553

FT /*tag= p

FT exon 1554..1751

FT /*tag= q

FT intron 1752..1815

FT exon /*tag= r
 FT 1816..1872
 FT /*tag= s
 FT 1873..1928
 FT intron /*tag= t
 FT 1929..2135
 FT exon /*tag= u
 FT 2136..2195
 FT intron /*tag= v
 FT 2196..2492
 FT exon /*tag= w
 PN WO9600290-A1.
 PD 04-JAN-1996.
 PF 15-JUN-1995; U07536.
 PR 24-JUN-1994; US-265534.
 PR 15-MAY-1995; US-441147.
 PA (NOVO) NOVO NORDISK BIOTECH INC.
 PA (NOVO) NOVO NORDISK AS.
 PI Aaslyng DA, Dalboge H, Schneider P, Xu F, Yaver DS;
 PI WPI; 96-068874/07.
 DR P-PSDB; R90722.
 PT DNA constructs for expression of Polyporus laccase enzymes - for use
 PT in e.g. lignin manipulation, juice mfr., phenol polymerisation and
 PT phenol resin prodn
 PS Claim 5; Page 62-65; 137pp; English.
 CC The sequence encodes laccase-LCC2 (pi 5.95) from Polyporus pinsitus
 CC (Trametes villosa). Polymerase chain reaction (PCR) amplification of
 CC P. pinsitus cDNA using primers 3331 (T15603) and 3332 (T15604) gives
 CC a 1500 bp fragment, which is joined to a signal peptide sequence
 CC from a 43-kDa cellulase using primer PH433 (T15605) and a pUC
 CC forward primer in PCR. Clones are expressed in Aspergillus oryzae,
 CC and a cDNA probe is obtained and used to screen a P. pinsitus
 CC genomic library in Escherichia coli DH5-alpha, giving plasmid
 CC pDSV19 (23GEN), with a 4-kb HindIII insert (NRRL B-21266).
 CC Screening also results in isolation of LCC1 (T15598) and LCC3-LCC5
 CC (T15600-T15602), which encode different laccases produced by P.
 CC pinsitus. The laccases may be used to polymerise lignin or
 CC lignosulphonates, to depolymerise Kraft pulp, to oxidise dyes or
 CC precursors, in hair dye compositions, or to polymerise or oxidise a
 CC phenolic or aniline compound. These new laccases are well-expressed
 CC in Aspergillus spp. (with vector integration in the genome), in
 CC contrast to previous basidiomycete laccases, which give low yields
 CC of recombinant enzyme. 547 A; 908 C; 793 G; 632 T;
 SQ Sequence 2880 BP; 547 A; 908 C; 793 G; 632 T;

alignment_scores:

Quality: 33.00 Length: 9
 Ratio: 4.125 Gaps: 0
 Percent Similarity: 88.889 Percent Identity: 66.667

alignment_block:

US-08-653-294-13 x T15599/rev ..

Align seg 1/1 to reverse of: T15599 from: 1 to: 2880

2 ArgLeuAlaIleArgLeuAspGluArg 10

||||: |||||||||||||||

1733 CGGGTCGAAGTTCGTCGATGACGG 1707

seq_name: N_Geneseq_36.V83943

seq_documentation_block:

ID V83943 standard; DNA; 11811 BP.

AC V83943;

DT 03-MAR-1999 (first entry)

DE Bacterial artificial chromosome (BAC)-F2 contig 3.

KW Yeast artificial chromosome; YAC; probe; eukaryotic chromosome;

KW neocentromere; replication; extra-chromosomal element; segregation;

KW cell division; artificial chromosome; gene therapy; BAC; transgenic;

KW human artificial chromosome; bacterial artificial chromosome; ss.

OS Synthetic.

PN WO9851790-A1.

```
PD 19-NOV-1998; AU0352.
PF 13-MAY-1998; AU008791.
PR 26-AUG-1997; AU-006784.
PR 13-MAY-1997; AU-006784.
PA (AMRA-) AMRAD OPERATIONS PTY LTD.
PI Cancilla MR, Choo K, Du Sart D;
DR WPI; 99-009773/01.
PT New isolated nucleic acid comprising neocentromere sequences from
PT eukaryotic chromosome - used to produce replicable, segregating
PT artificial chromosomes that can carry large amounts of DNA for gene
PT therapy.
PS Claim 10; Page 195-203; 540pp; English.
CC The present sequence represents a bacterial artificial chromosome (BAC)
CC contig, and exemplifies the invention. The specification describes
CC nucleic acid sequences derived from a eukaryotic chromosome, including a
CC neocentromere or its functional derivative or hybrid, that are able, in
CC a compatible cell, of replicating, acting as extra-chromosomal element
CC and segregating during cell division. The sequences can be used to
CC construct artificial chromosomes for use in gene therapy comprising a
CC replicable, segregating nucleic acid that confers a specific phenotype
CC on cells. Human artificial chromosomes can propagate in human cells and
CC carry large amounts of DNA (e.g. therapeutic genes), and, being
CC extra-chromosomal, they are not mutagenic. The artificial chromosomes
CC are also useful for generation of transgenic plants and animals, in
CC production of proteins and to make diagnostic reagents, e.g. for
CC expression of cytokines, receptors and growth factors, or to increase
CC the copy number of a gene in a cell. The constructs may also be
CC used for functional and structural analysis of chromosomes.
SQ Sequence 11811 BP; 3014 A; 2459 C; 2433 G; 3905 T;

alignment_scores:
  Quality: 33.00      Length: 10
  Ratio: 3.667       Gaps: 0
Percent Similarity: 90.000 Percent Identity: 50.000

alignment_block:
US-08-653-294-13 x V83943/rev ..
  Align seg 1/1 to reverse of: V83943 from: 1 to: 11811

      1 TyrArgLeuAlaIleArgLeuAspGluArg 10
      |||||:||||:||||:||||:||||:||||:
4851 TACAGAAATTCAGTCAGGAGGATCAAAAA 4822

seq_name: N_Geneseq_36:T07040
seq_documentation_block:
ID T07040;
AC T07040 standard; DNA; 612 BP.
DT 03-JUL-1996 (first entry)
DE Immunogen DNA from n-(ABCEDE) hepatitis virus.
KW Non-A, non-B, non-C, non-D, non-E hepatitis virus; n-(ABCEDE);
KW immunogen; antibody; vaccine; phage library; ds.
OS Non-A, non-B, non-C, non-D, non-E hepatitis virus JFA clone 4B11.
PN WO9532290-A2.
PD 30-NOV-1995.
PF 17-MAY-1995; U05980.
PR 20-MAY-1994; US-246986.
PA (GENE-) GENELABS TECHNOLOGIES INC.
PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.
PI Kim JP, Purcell RH;
DR WPI; 96-020585/02.
PT New non-A, -B, -C, -D and -E (n-(ABCEDE)) hepatitis DNA libraries -
PT used to develop prods. for the detection, diagnosis, prevention and
PT treatment of n-(ABCEDE) hepatitis.
PS Disclosure; Page 113; 165pp; English.
CC The sequence represents clone 4B11 which encodes an immunogenic
CC polypeptide associated with non-A, non-B, non-C, non-D, non-E
CC (n-(ABCEDE)) hepatitis virus infection, and is obtained by
CC preparing a phage from a phage lambda gtl1 library of JFA DNA
CC (ATCC 75271) n-(ABCEDE) hepatitis serum, plating to form plaques,
CC and screening the phage plaques for the production of polypeptides
```

```
CC immunoreactive with n-(ABCEDE) serum. Inserted sequences in gtl1
CC are expressed as beta-galactosidase fusion proteins. Clone 4B11
CC has 3 multiple insert sequences; each insert can be fractionated
CC into discrete sequences by restriction enzymes, or portions of the
CC inserts can be PCR amplified by sequence specific primers. Each
CC resulting individual region can be subcloned and immunoscreened.
CC This allows identification of specific regions responsible for
CC immunoreactivity. n-(ABCEDE) hepatitis virus polypeptides can be
CC used for the production or detection of antibodies, and in
CC vaccines. The antibodies can be used for detection, diagnosis and
CC in passive immunotherapy. The DNA can be used in detection and
CC diagnosis, and as hybridisation probes for identification of
CC further n-(ABCEDE) hepatitis virus coding sequences. Culture systems
CC producing the n-(ABCEDE) polypeptides can be used in screening
CC studies.
SQ Sequence 612 BP; 153 A; 171 C; 161 G; 127 T;

alignment_scores:
  Quality: 32.00      Length: 9
  Ratio: 3.556       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 66.667

alignment_block:
US-08-653-294-13 x T07040/rev ..
  Align seg 1/1 to reverse of: T07040 from: 1 to: 612

      2 ArgLeuAlaIleArgLeuAspGluArg 10
      |||||:||||:||||:||||:||||:
49 CGCCTGGCGCTCGAAGTCGATGACGCG 23

seq_name: N_Geneseq_36:T34189
seq_documentation_block:
ID T34189 standard; DNA; 972 BP.
AC T34189;
DT 25-SEP-1996 (first entry)
DE PCpD coding sequence.
KW Pentachlorophenol breakdown pathway: PCpC; PCpA; PCpB; environment;
KW tetrachloro-p-hydroquinone reductase; PCp-degrading enzyme complex;
KW Flavobacterium; PCp; 2,3,5,6-tetrachloro-p-hydroquinone; TeCH;
KW glutathione; 2,3,6-trichloro-p-hydroquinone; TrCH; food chain;
KW 2,6-dichloro-p-hydroquinone; wood preserving industry; fungicide;
KW pesticide; herbicide; disinfectant; ds.
OS Flavobacterium sp. Strain ATCC 39723.
FH Key Location/Qualifiers
FT misc_difference 754
FT /note= "Residue not given in the specification, included
FT to maintain open reading frame and to encode the
FT amino acid given in the specification at this
FT position (Glu)"
PN US5512478-A.
PD 30-APR-1996.
PF 23-MAR-1992; 856015.
PR 23-MAR-1992; US-856015.
PR 13-JUL-1992; US-914282.
PR 18-JUL-1994; US-276887.
PA (DAH-) IDAHO RES FOUND INC.
PI Lange CC, Orser CS, Xun L;
DR WPI; 96-229872/23.
DR P-PSDB; R99487.
PT Flavobacterium sp. penta:chloro:phenol breakdown pathway genes and
PT enzymes - useful in the bio-remediation and dechlorination of PCP
PT contg. matter
PS Disclosure; Columns 61-64; 52pp; English.
CC The sequences given in T34189-90 encode proteins from the
CC pentachlorophenol (PCp) breakdown pathway, designated PCpD and
CC PCpR. The PCpD gene was found to code for a 323 amino acid
CC polypeptide, mol. wt. 35942 daltons. Based on multiple sequence
CC alignments, PCpD belongs to the family of class 1 dioxygenase electron
CC transport proteins. PCpD was predicted to have three structural
```

CC domains, one involved in binding flavin mononucleotide (FMN), the second
 CC responsible for binding reduced NADPH and the third forming a plant-
 CC ferredoxin-type (3Fe-2S) centre. PcpD is classified as an iron
 CC sulphur flavoprotein-oxidoreductase, and has been designated the PCP
 CC 4-monooxygenase reductase. It is transcribed as a dicistronic message
 CC with PcpB (see also T34184). The pcpR open reading frame encodes a 303
 CC amino acid protein with a mol. wt. of 33549 daltons, which possesses a
 CC helix-turn-helix motif in its N-terminal portion. PcpR is thought to
 CC activate the transcription of pcpBD and pcpA. In combination with pcpA,
 CC pcpB and pcpC, these enzymes catalyse the initial steps of PCP breakdown.
 CC These enzymes can specifically be used in the breakdown of PCP
 CC containing matter which persists in the environment and in food chains
 CC after its use in the wood preserving industry as a fungicide and
 CC pesticide, and in products such as herbicides and disinfectants.
 SQ Sequence 972 BP; 181 A; 310 C; 313 G; 168 T;

alignment_scores:
 Quality: 32.00 Length: 8
 Ratio: 4.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 75.000

alignment_block:

US-08-653-294-13 x T34189/rev ..
 Align seg 1/1 to reverse of: T34189 from: 1 to: 972

3 LeuAlaIleArgLeuAspGluArg 10
 |||||:||||:|||||
 192 CTCGCCCTGCGGGTTCGACGACGA 169

seq_name: N_Geneseq_36:V84117

seq_documentation_block:

ID V84117 standard; DNA; 1038 BP.
 AC V84117;
 DT 15-MAR-1999 (first entry)
 DE Pseudomonas aeruginosa heptosyl transferase II waaF gene.
 KW waaF gene; rfaF gene; lipopolysaccharide; infection; therapy;
 KW diagnosis; vaccine; heptosyl transferase II; ss.
 OS Pseudomonas aeruginosa strain PAO1.
 PN W09850557-Al.
 PD 12-NOV-1998.
 PF 01-MAY-1998; CA0395.
 PR 09-MAY-1997; US-046149.
 PR 02-MAY-1997; US-045418.
 PA (UYGU-) UNIV GUELPH.
 PI Burrows LL, De Kievit TR, Lam JS, Matewish M, Walsh A;
 DR WPI; 99-034725/03.
 DR P-PSDB; W88211.
 PT Isolated P. aeruginosa waaC, waaG, waaF and waaP gene cluster -
 PT useful in the diagnosis or treatment of P. aeruginosa infections
 PS Claim 2; Fig 6; 61pp; English.
 CC This is the nucleotide sequence of the waaF gene (rfaF gene) of
 CC the waa gene cluster of Pseudomonas aeruginosa PAO1. It codes for
 CC waaF (see W82211), a heptosyl transferase II that adds the second
 CC heptose residue onto the core oligosaccharide in the biosynthesis
 CC of the lipopolysaccharide inner core. The four waa genes
 CC of P. aeruginosa (see V84116-19) are arranged contiguously in an
 CC operon with the gene order waaF, waaC, waaG and waaP. The
 CC functions of the encoded proteins (see W82210-13) were tested by
 CC complementation of specific Salmonella typhimurium mutants, and
 CC knockout mutations of the genes in P. aeruginosa. The waa nucleic
 CC acids or proteins can be used to diagnose a bacterial, especially a
 CC P. aeruginosa, infection in an animal. They can further be used to
 CC screen for compounds that affect core lipopolysaccharide biosynthesis
 CC or assembly. A claimed method of treating or preventing a bacterial
 CC infection comprises administering an agent that inhibits a waa gene
 CC to an animal. A claimed vaccine for treating a bacterial infection
 CC includes one of WaaP, WaaC, WaaF and WaaG.
 SQ Sequence 1038 BP; 148 A; 348 C; 364 G; 178 T;

alignment_scores:
 Quality: 32.00 Length: 8
 Ratio: 4.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 75.000

alignment_block:

US-08-653-294-13 x V84117 ..
 Align seg 1/1 to: V84117 from: 1 to: 1038

3 LeuAlaIleArgLeuAspGluArg 10
 |||||:||||:|||||
 630 CTGCTGTTCTGGCTCGAAGAACGA 653

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:45 ; Search time 209.03 Seconds
(without alignments)
3.980 Million cell updates/sec

Title: US-08-653-294-18
Perfect score: 58
Sequence: 1 YRLAIRLLRLY 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 225878 seqs, 59334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database :

SPTREMBL_12:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phase:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	37	63.8	259	11 Q60503	Q60503 cricetus
2	37	63.8	321	5 Q45795	Q45795 caenorhabdi
3	36	62.1	141	2 Q84514	Q84514 chlamydia t
4	36	62.1	243	2 Q85853	Q85853 sphingomona
5	36	62.1	309	2 Q86347	Q86347 mycobacteri
6	36	62.1	725	5 Q16382	Q16382 caenorhabdi
7	36	62.1	795	5 Q22063	Q22063 caenorhabdi
8	36	62.1	1847	5 P91495	P91495 caenorhabdi
9	35	62.1	1899	10 Q9XEG1	Q9XEG1 gossypium h
10	35	60.3	142	2 Q9Z759	Q9Z759 chlamydia p
11	35	60.3	325	2 Q9XB05	Q9XB05 myxococcus
12	35	60.3	350	5 Q45803	Q45803 caenorhabdi
13	35	60.3	683	5 Q9Y136	Q9Y136 drosophila
14	35	60.3	1283	5 Q24393	Q24393 drosophila
15	35	60.3	1605	5 Q96446	Q96446 vairimorpha
16	34	58.6	61	12 Q9WHV4	Q9WHV4 squash yell
17	34	58.6	111	2 P70780	P70780 anabaena sp
18	34	58.6	152	2 Q9WYM7	Q9WYM7 thermotoga
19	34	58.6	261	2 Q9Z671	Q9Z671 zymomonas m
20	34	58.6	280	2 Q35043	Q35043 bacillus su

21	34	58.6	285	5 P91068	P91068 caenorhabdi
22	34	58.6	377	4 Q60704	Q60704 homo sapien
23	34	58.6	735	5 Q76912	Q76912 drosophila
24	34	58.6	777	4 Q43162	Q43162 homo sapien
25	34	58.6	814	10 Q64487	Q64487 arabidopsis
26	34	58.6	1712	5 Q96160	Q96160 plasmodium
27	33	56.9	35	5 Q61237	Q61237 onchocerca
28	33	56.9	59	7 Q78094	Q78094 homo sapien
29	33	56.9	133	2 Q92903	Q92903 chlamydia p
30	33	56.9	168	1 Q28893	Q28893 archaeglob
31	33	56.9	184	3 Q13954	Q13954 schizosacch
32	33	56.9	194	5 Q21212	Q21212 caenorhabdi
33	33	56.9	260	12 Q9YW71	Q9YW71 melanoplus
34	33	56.9	313	2 P97213	P97213 clostridium
35	33	56.9	326	2 Q9ZJ72	Q9ZJ72 helicobacte
36	33	56.9	334	2 Q52631	Q52631 clostridium
37	33	56.9	345	10 Q24513	Q24513 arabidopsis
38	33	56.9	352	2 Q9X816	Q9X816 streptomyce
39	33	56.9	427	5 Q22310	Q22310 caenorhabdi
40	33	56.9	452	2 Q59958	Q59958 streptococc
41	33	56.9	453	2 Q83072	Q83072 treponema p
42	33	56.9	458	5 Q19489	Q19489 caenorhabdi
43	33	56.9	532	2 P72007	P72007 mycobacteri
44	33	56.9	576	2 Q84579	Q84579 chlamydia t
45	33	56.9	650	2 Q52818	Q52818 amycolatops

ALIGNMENTS

RESULT 1
Q60503 PRELIMINARY; PRT; 259 AA.
AC Q60503;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-NOV-1998 (TREMBlrel. 08, Last annotation update)
DE P-GLYCOPROTEIN (FRAGMENT).

GN Cricetulus griseus (Chinese hamster).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Cricetinae; Cricetulus.
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE-OVARY;
RX MEDLINE; 89261726.
RA NG W.F., SARANGI F., ZASTAWNY R.L., VEINOT-DREBOT L., LING V.;
RT "Identification of members of the P-glycoprotein multigene family."
RL Mol. Cell. Biol. 9:1224-1232(1989).
DR EMBL; M25792; AAA53439.1; JOINED.
DR EMBL; M25789; AAA53439.1; JOINED.
DR EMBL; M25790; AAA53439.1; JOINED.
DR EMBL; M25791; AAA53439.1; JOINED.
DR PFAM; PF00005; ABC_tran; 1.
FT NON_TER 1
SQ SEQUENCE 259 AA; 28543 MW; 021E9AF3 CRC32;

Query Match 63.8%; Score 37; DB 11; Length 259;
Best Local Similarity 80.0%; Pred. No. 18;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLAIIRLLR 11
| | | | | | | |
Db 162 RLAIIRLLR 171

RESULT 2
Q45795 PRELIMINARY; PRT; 321 AA.
ID Q45795;
AC Q45795;
DT 01-JUN-1998 (TREMBlrel. 06, Created)
DT 01-JUN-1998 (TREMBlrel. 06, Last sequence update)

DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
 GN T19C9.4
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Secernentes; Rhabditia; Rhabditida;
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA MATTHEWS L.;
 RL Submitted (MAR-1997) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 94150718.
 RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,
 RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
 RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
 RA GARDNER A., GREEN P., HAWKINS T., HILLIER J., JIER M., JOHNSTON L.,
 RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
 RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
 RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
 RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
 RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
 RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 RT elegans.";
 RL Nature 368:32-38(1994).
 DR EMBL; 292972; CAB07489.1; -
 DR PFAM; PF01604; 7tm5.1;
 SQ SEQUENCE 321 AA; 36504 MW; 70031B52 CRC32;

Query Match 63.8%; Score 37; DB 5; Length 321;
 Best Local Similarity 88.9%; Pred. No. 22;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRRL 9
 DB 298 YRAAIRRL 306
 ||| |||||

RESULT 3
 ID 084514 PRELIMINARY; PRT; 141 AA.
 AC 084514;
 DT 01-NOV-1998 (TREMBlrel. 08, Created)
 DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)
 DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
 DE 50S RIBOSOMAL PROTEIN L17.
 GN RL17.
 OS Chlamydia trachomatis.
 OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=D/UN-3/CX;
 RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L.,
 RA MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V.,
 RA DAVIS R.W.;
 RT "Genome Sequence of an Obligate Intracellular Pathogen of Humans:
 RT Chlamydia trachomatis.";
 RL Science 0:0-0(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=D/UN-3/CX;
 RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L.,
 RA MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V.,
 RA DAVIS R.W.;
 RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
 CC -!- SIMILARITY: BELONGS TO THE L17F FAMILY OF RIBOSOMAL PROTEINS.
 DR EMBL; AAC01323; AAC68107.1;
 DR PROSITE; PS01167; RIBOSOMAL_L17; 1.
 DR PFAM; PF01196; Ribosomal_L17; 1.
 KW Ribosomal protein.
 SQ SEQUENCE 141 AA; 16152 MW; 2570FF77 CRC32;

Query Match 62.1%; Score 36; DB 2; Length 141;
 Best Local Similarity 54.5%; Pred. No. 15;
 Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLAIRRL 12
 DB 64 RLAVGLMVR 74
 ||| |::|

RESULT 4
 ID 085853 PRELIMINARY; PRT; 243 AA.
 AC 085853;
 DT 01-NOV-1998 (TREMBlrel. 08, Created)
 DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)
 DT 01-NOV-1998 (TREMBlrel. 08, Last annotation update)
 DE HYPOTHETICAL 26.5 KD PROTEIN.
 OS Spingomonas aromaticivorans.
 OG Plasmid pNL1.
 OC Bacteria; Proteobacteria; alpha subdivision; Zymomonas group;
 OC Spingomonas.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=F199;
 RA ROMINE M.F., STILLWELL L.C., WONG K.-K., THURSTON S.J., SISK E.C.,
 RA SENSEN C.W., GAASTERLAND T., SAFFER J.D., FREDRICKSON J.K.;
 RT "Complete sequence of a 184 kb catabolic plasmid from Spingomonas
 RT aromaticivorans strain F199.";
 RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AF079317; AAO03868.1; -
 KW Hypothetical protein; Plasmid.
 SQ SEQUENCE 243 AA; 26455 MW; 40CDFBF4 CRC32;

Query Match 62.1%; Score 36; DB 2; Length 243;
 Best Local Similarity 63.6%; Pred. No. 26;
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIRRL 12
 DB 5 RLQRRVTIR 15
 ||| |||::|

RESULT 5
 ID 086347 PRELIMINARY; PRT; 309 AA.
 AC 086347;
 DT 01-NOV-1998 (TREMBlrel. 08, Created)
 DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)
 DT 01-NOV-1999 (TREMBlrel. 12, Last annotation update)
 DE HYPOTHETICAL 33.5 KD PROTEIN.
 GN RV2776C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=H37RV;
 RX MEDLINE; 98295987.
 RA COLE S.T., BROSCHE R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
 RA GORDON S.V., EIGLMEIER K., GAS S., BARRY III C.E., TEKAIA F.,
 RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
 RA DAVIES R., DEVLIN K., FELTWELL T., GENTLES S., HAMLIN N., HOLROYD S.,
 RA HORNSBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,
 RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAN M.A., ROGERS J.,
 RA RUTTER S., SEEGER K., SKELTON S., SQUARES S., SQUARES R., SULSTON J.E.,
 RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 RT complete genome sequence.";
 RL Nature 393:537-544(1998).
 RN [2]

RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RA PARKHILL J.;

RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL: AL008967; CAA15591.1; -

DR HSSP: P33164; 2PIA.

DR PROSITE: PS00197; 2FE2S_FERREDOXIN; 1.

DR PFAM: PF00111; Ier2; 1.

DR PFAM: PF00175; oxidored_fad; 1.

KW Hypothetical protein; Iron-sulfur.

SQ SEQUENCE 309 AA; 33517 MW; B152B590 CRC32;

Query Match 62.1%; Score 36; DB 2; Length 309;

Best Local Similarity 87.5%; Pred. No. 33;

Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRRI 8

DB 65 YRLAIRRI 72

RESULT 6

QY 016382

AC 016382 PRELIMINARY; PRT; 725 AA.

DT 01-JAN-1998 (TREMREL. 05, Created)

DT 01-JAN-1998 (TREMREL. 05, Last sequence update)

DT 01-NOV-1998 (TREMREL. 08, Last annotation update)

DE K12B6.2 PROTEIN.

GN K12B6.2.

OS Caenorhabditis elegans.

OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;

OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.

[1]

RP SEQUENCE FROM N.A.

RC STRAIN=BRISTOL N2;

RX MEDLINE; 94150718.

RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,

RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,

RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,

RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,

RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,

RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,

RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,

RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULTON J.,

RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,

RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.,

RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.

elegans.";

RL Nature 368:32-38(1994).

[2]

RP SEQUENCE FROM N.A.

RC STRAIN=BRISTOL N2;

RA JONES K., WOHLDMANN P.;

RL Submitted (AUG-1997) to the EMBL/GenBank/DBJ databases.

[3]

RP SEQUENCE FROM N.A.

RC STRAIN=BRISTOL N2;

RA WATERSTON R.;

RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.

DR EMBL: AF016442; AAF65917.1; -

SQ SEQUENCE 725 AA; 84926 MW; A4847D75 CRC32;

Query Match 62.1%; Score 36; DB 5; Length 725;

Best Local Similarity 58.3%; Pred. No. 77;

Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRRILLY 12

DB 594 FFAIRRILVLY 605

RESULT 7

QY 022063

AC 022063 PRELIMINARY; PRT; 795 AA.

DT 01-NOV-1996 (TREMREL. 01, Created)

DT 01-MAY-1999 (TREMREL. 10, Last sequence update)

DT 01-NOV-1999 (TREMREL. 12, Last annotation update)

DE T01C3.10 PROTEIN.

GN T01C3.10.

OS Caenorhabditis elegans.

OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;

OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.

[1]

RP SEQUENCE FROM N.A.

RA WILD A.;

RL Submitted (AUG-1996) to the EMBL/GenBank/DBJ databases.

DR EMBL: Z81061; CAB02939.1; -

DR EMBL: Z78413; CAB02939.1; JOINED.

DR EMBL: Z78413; CAB01667.1; -

DR EMBL: Z81061; CAB01667.1; JOINED.

DR HSSP: P19491; IGR2.

DR PFAM: PF00060; lig_chan; 1.

SQ SEQUENCE 795 AA; 89703 MW; DD722166 CRC32;

Query Match 62.1%; Score 36; DB 5; Length 795;

Best Local Similarity 50.0%; Pred. No. 85;

Matches 6; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRRILLY 12

DB 6 YRTSLRLATRY 17

RESULT 8

QY 022063

AC 022063 PRELIMINARY; PRT; 1847 AA.

DT 01-MAY-1997 (TREMREL. 03, Created)

DT 01-MAY-1997 (TREMREL. 03, Last sequence update)

DT 01-NOV-1998 (TREMREL. 08, Last annotation update)

DE SIMILARITY TO RAT INTEGRAL MEMBRANE GLYCOPROTEIN GP120 PRECURSOR.

GN T23H2.1.

OS Caenorhabditis elegans.

OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;

OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.

[1]

RP SEQUENCE FROM N.A.

RC STRAIN=BRISTOL N2;

RX MEDLINE; 94150718.

RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,

RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,

RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,

RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,

RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,

RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,

RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,

RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULTON J.,

RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,

RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.,

RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.

elegans.";

RL Nature 368:32-38(1994).

[2]

RP SEQUENCE FROM N.A.

RC STRAIN=BRISTOL N2;

RA WATSON P., BRADSHAW H.;

RL Submitted (JAN-1997) to the EMBL/GenBank/DBJ databases.

DR EMBL: U80033; AAC48199.1; -

SQ SEQUENCE 1847 AA; 199637 MW; B0A25E0F CRC32;

Query Match 62.1%; Score 36; DB 5; Length 1847;
 Best Local Similarity 58.3%; Pred. No. 1.9e+02;
 Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRRLLRY 12
 DB 19 YRLNPRVLLPY 30
 ||| : ||| |

RESULT 9
 QXEG1 PRELIMINARY; PRT: 1899 AA.
 ID QXEG1
 AC QXEG1;
 DT 01-NOV-1999 (TRENBLrel. 12, Created)
 DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE PUTATIVE CALLOSE SYNTHASE CATALYTIC SUBUNIT.
 GN CFL1.
 OS Gossypium hirsutum (Upland cotton).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC eukaryotes; Spermatophyta; Magnoliophyta; eudicotyledons;
 OC core eudicots; Rosidae; eurosids II; Malvales; Malvaceae; Gossypium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-CV. TEXAS MARKER-1; TISSUE=PRIMARY-STAGE COTTON FIBER;
 RA CUI X., SHIN H., BROWN R.M.;
 RT "Cotton CFL1 gene shows homology to the yeast beta-1,3-glucan synthase
 subunit FKS1."
 RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF085717; RAD25952.1; -
 SQ SEQUENCE 1899 AA; 218627 MW; E695145F CRC32;

Query Match 62.1%; Score 36; DB 10; Length 1899;
 Best Local Similarity 63.6%; Pred. No. 2e+02;
 Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIRRLLRY 12
 DB 554 RLAVSRIFLRF 564
 ||| : ||| |

RESULT 10
 QZ7S9 PRELIMINARY; PRT: 142 AA.
 ID QZ7S9;
 AC QZ7S9;
 DT 01-MAY-1999 (TRENBLrel. 10, Created)
 DT 01-MAY-1999 (TRENBLrel. 10, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE 50S RIBOSOMAL PROTEIN L17.
 GN RL17.
 OS Chlamydia pneumoniae.
 OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydiaophila.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-CWL029;
 RA KALMAN S., MITCHELL W., MARATHE R., LAMMEL C., FAN J., OLINGER L.,
 RA GRIMWOOD J., DAVIS R.W., STEPHENS R.S.;
 RT "Comparative Genomes of Chlamydia pneumoniae and C. trachomatis."
 RL Submitted (DEC-1998) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: BELONGS TO THE L17P FAMILY OF RIBOSOMAL PROTEINS.
 DR EMBL: AE001647; RAD16764.1; -
 DR PROSITE: PS01167; RIBOSOMAL_L17; 1.
 KW Ribosomal protein.
 SQ SEQUENCE 142 AA; 16400 MW; 4839EC84 CRC32;

Query Match 60.3%; Score 35; DB 2; Length 142;
 Best Local Similarity 54.5%; Pred. No. 24;
 Matches 6; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLAIRRLLRY 12
 ||| : ||| |

Db 64 RIAIGRLMRY 74
 RESULT 11
 Q9XB05 PRELIMINARY; PRT: 325 AA.
 ID Q9XB05
 AC Q9XB05;
 DT 01-NOV-1999 (TRENBLrel. 12, Created)
 DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE MEMBRANE ASSOCIATED PROTEIN.
 GN TAD.
 OS Myxococcus xanthus.
 OC Bacteria; Proteobacteria; delta subdivision; Myxobacteria;
 OC Myxococcales; Cystobacterineae; Myxococcaceae; Myxococcus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-ER-15;
 RA PATAN Y., ORR E., RON E.2., ROSENBERG E.;
 RT "Genetic and functional analysis of genes required for the post-
 modification of the polyketide antibiotic TA of Myxococcus xanthus."
 RL Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AJ132503; CAB46503.1; -
 SQ SEQUENCE 325 AA; 35985 MW; 4CC64E85 CRC32;

Query Match 60.3%; Score 35; DB 2; Length 325;
 Best Local Similarity 58.3%; Pred. No. 54;
 Matches 7; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRRLLRY 12
 DB 202 YRLTVDREPLY 213
 ||| : ||| |

RESULT 12
 Q45803 PRELIMINARY; PRT: 350 AA.
 ID Q45803
 AC Q45803;
 DT 01-JUN-1998 (TRENBLrel. 06, Created)
 DT 01-JUN-1998 (TRENBLrel. 06, Last sequence update)
 DT 01-MAY-1999 (TRENBLrel. 10, Last annotation update)
 DE T23D5.1 PROTEIN.
 GN T23D5.1.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA LLOYD C., WILKINSON J.;
 RL Submitted (NOV-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE: 94150718.
 RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,
 RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
 RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
 RA GARDNER A., GREEN P., HARKINS T., HILLIER L., JIER M., JOHNSTON L.,
 RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
 RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
 RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
 RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
 RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
 RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 elegans."
 RL Nature 368:32-38(1994).
 DR EMBL: Z82051; CAB04815.1; -
 DR PFWA: PF01461; 7tm.4; 1.
 SQ SEQUENCE 350 AA; 40291 MW; A9117B4D CRC32;

Query Match 60.3%; Score 35; DB 5; Length 350;
 ||| : ||| |

Best Local Similarity 66.7%; Pred. No. 58;
Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 1 YRLAIRRIL 9
|:|:|:|:|:
DB 307 YRLAVRKIV 315

RESULT 13
QY136 PRELIMINARY; PRT; 683 AA.
ID QY136
AC QY136
DT 01-NOV-1999 (TREMREL. 12, Created)
DT 01-NOV-1999 (TREMREL. 12, Last sequence update)
DT 01-NOV-1999 (TREMREL. 12, Last annotation update)
DE BCDNA.GH07188.
DE BCDNA.GH07188.
GN Drosophila melanogaster (Fruit fly).
OS Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A.
RA RUBIN G.M., WAN K.H., HARVEY D., LEWIS S.E., BROKSTEIN P., TSANG G.,
RA AGRAYANI A., ARCAINA T.T., BAXTER E., BLAZEJ R.G., BUTENHOFF C.,
RA CHAMPE M., CHAVEZ C., CHEW M., DOYLE C.M., FARFAN D.E., FRISSE E.,
RA GALLE R., GEORGE R.A., HARRIS N.L., HOSKINS R.A., EVANS-HOLM M.,
RA HOUSTON K.A., HUMMASTI S.R., KIM E., LI P., MOSHREFI M., PACLEB J.M.,
RA PARK S., SEQUEIRA A., SETHI H., SNIR E., SVIRSKAS R.R., WEINBURG T.,
RA CELNIKER S.E.
RT "Full length Drosophila melanogaster cDNA sequence."
RL Submitted (APR-1999) to the EMBL/Genbank/DBSJ databases.
DR EMBL; AF145636; AAD38611.1;
SQ SEQUENCE 683 AA; 79171 MW; 72C620E0 CRC32;

Query Match 60.3%; Score 35; DB 5; Length 683;
Best Local Similarity 66.7%; Pred. No. 1.le+02;
Matches 8; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRRILLY 12
|:|:|:|:|:
DB 206 YRNNIQRILQRY 217

RESULT 14
Q24393 PRELIMINARY; PRT; 1283 AA.
ID Q24393
AC Q24393
DT 01-NOV-1996 (TREMREL. 01, Created)
DT 01-NOV-1996 (TREMREL. 01, Last sequence update)
DT 01-NOV-1999 (TREMREL. 12, Last annotation update)
DE P-GLYCOPROTEIN/MULTIDRUG RESISTANCE PROTEIN.
GN MDR50.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-CANTON S;
RX MEDLINE: 94010914.
RA GERRARD B., STEWART C., DEAN M.;
RT "Analysis of Mdr50: a Drosophila p-glycoprotein/multidrug resistance gene homolog."
RL Genomics 17:83-88(1993).
DR EMBL; L07065; AAL16186.1;
DR FLYBASE: FBgn010241; Mdr50.
DR PROSITE: PS00211; ABC_TRANSPORTER; 1.
DR PFAM: PF00664; ABC_membrane; 2.
DR PFAM: PF00005; ABC_tran; 2.
KW ATP-binding; Transport.
SQ SEQUENCE 1283 AA; 142538 MW; 65012909 CRC32;

Query Match 60.3%; Score 35; DB 5; Length 1283;
Best Local Similarity 70.0%; Pred. No. 2.1e+02;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLAIIRILLR 11
|:|:|:|:|:
DB 547 RLAIIRALIR 556

RESULT 15
O96446 PRELIMINARY; PRT; 1605 AA.
ID O96446
AC O96446;
DT 01-MAY-1999 (TREMREL. 10, Created)
DT 01-MAY-1999 (TREMREL. 10, Last sequence update)
DT 01-MAY-1999 (TREMREL. 10, Last annotation update)
DE LARGEST SUBUNIT OF RNA POLYMERASE II.
GN RPBI.
OS Vairimorpha necatrix.
OC Eukaryota; Microsporidia; Burenellidae; Vairimorpha.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 99110933.
RA HIRT R.P., LOGSDON J.M. JR., HEALY B., DOREY M.W., DOOLITTLE W.F.,
RA EMBLEY T.M.;
RT "Microsporidia are related to Fungi: evidence from the largest subunit of RNA polymerase II and other proteins."
RL Proc. Natl. Acad. Sci. U.S.A. 96:580-585(1999).
DR EMBL; AF060234; AAD12604.1;
SQ SEQUENCE 1605 AA; 180946 MW; 2D013184 CRC32;

Query Match 60.3%; Score 35; DB 5; Length 1605;
Best Local Similarity 58.3%; Pred. No. 2.6e+02;
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRRILLRY 12
|:|:|:|:|:
DB 1009 YNLSIKRILNEY 1020

Search completed: February 8, 2000, 13:17:46
Job time: 32495 sec

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OM of: US-08-653-294-18 to: GenEmbl.* out_format : pfs

Date: Feb 8, 2000 4:42 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

-MODEL=frame+p2n.model -DEV=xlp
-O=Cgml_1/USPTO.spool/US08653294/runat_04022000.160701.15779/app_query.fasta.1
-DB=GenEmbl -QFMT=fastap -SUFFIX=rge -GAPOP=12.000 -GAPEXT=4.000
-MINMATCH=0.100 -LOOPEXT=0.000 -LOOPEXT=0.000 -GAPOP=4.500
-GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
-FGAPOP=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DELOP=6.000
-DELEXT=7.000 -START=1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOALIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
-OUTFMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=1000000 -USER=US08653294
-NCPU=6 -ICPU=3 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-08-653-294-18

Query length: 12

Database: GenEmbl.*

Database sequences: 821193

Database length: 1518192014

Search time (sec): 11370.480000

score_list:

Sequence	Strd	Orig	ZScore	Escore	Len	Documentation
gb_hg7:AC013772	-	47.00	110.79	378.51	142796	..
gb_hg7:HS076555C5	+	45.00	96.17	2.5e+03	AC013772	Homo sapiens clone
gb_hg5:AC011145	+	44.00	98.46	1.8e+03	AC0121956	Homo sapiens clone
gb_hg4:AC012150	+	42.00	103.73	936.38	AC011145	Homo sapiens clone
gb_in1:DMR3A3	+	41.00	132.31	23.96	37459	AC012150
gb_pr2:MH01AFHOM	+	41.00	132.26	24.12	1056	V00238
gb_hg7:AC017966	+	41.00	112.98	286.02	8849	Three Drosophila melanogaster
gb_hg5:AC013927	+	41.00	109.65	438.30	AC017966	Drosophila melanogaster
gb_bal:MLC82533	+	41.00	99.24	1.7e+03	AC013927	Drosophila melanogaster
gb_bal:000017	+	41.00	98.82	1.8e+03	AC003510	Mycobacterium leprae
gb_hg6:AC007807	+	41.00	88.56	6.5e+03	AC007807	Drosophila melanogaster
gb_hg4:AC010564	+	41.00	87.68	7.3e+03	AC010564	Drosophila melanogaster
gb_hg2:AC005047	+	41.00	86.47	8.2e+03	AC005047	Homo sapiens clone
gb_pr4:AC007687	+	41.00	86.46	8.6e+03	AC007687	Homo sapiens clone
gb_hg1:HS392M18	+	41.00	83.99	1.2e+04	AC0121897	Homo sapiens clone
gb_in2:AC005847	+	41.00	82.03	1.5e+04	AC005847	Drosophila melanogaster
gb_hg3:AC008835	+	40.00	105.61	735.67	AC008835	Homo sapiens chromosome
gb_in1:CEFI14B6	+	40.00	98.28	1.9e+03	AC010564	Drosophila melanogaster
gb_pr1:SPBC725	+	40.00	95.93	2.5e+03	AC034352	S.pombe chromosome
gb_pr1:AB028605	+	40.00	95.90	2.6e+03	AB028605	Arabidopsis thaliana
gb_hg5:AC015514	+	40.00	89.30	6.0e+03	AC015514	Homo sapiens clone
gb_hg6:AC015746	+	40.00	87.95	7.1e+03	AC015746	Homo sapiens clone
gb_hg5:AC015908	+	40.00	81.92	1.5e+04	AC015908	Homo sapiens chromosome
gb_hg1:CEY47H10	+	40.00	77.28	2.8e+04	AC015908	Homo sapiens chromosome
gb_pr2:CN0191TC	+	39.00	136.00	14.94	AC012072	Botrytis cinerea strain
gb_ro:MMU16741	+	39.00	120.31	111.62	U16741	Mus musculus capping protein
gb_bal:AF025541	+	39.00	120.30	111.77	AF025541	Mycobacterium tuberculosis
gb_pr1:AB007931	+	39.00	107.23	597.75	AB007931	Mycobacterium tuberculosis
gb_hg3:AC009605	+	39.00	92.57	3.9e+03	AC009605	Leishmania major
gb_in1:CELF45E1	+	39.00	92.05	4.2e+03	AC009605	Leishmania major
gb_bal:HSJ11266	+	39.00	91.93	4.3e+03	U28732	Caenorhabditis elegans
gb_hg5:AC012873	+	39.00	91.41	4.5e+03	U28732	Caenorhabditis elegans
gb_hg5:AC013205	+	39.00	88.52	6.6e+03	U28732	Caenorhabditis elegans
gb_pr3:AC003365	+	39.00	84.65	1.1e+04	AC012873	Drosophila melanogaster
gb_hg2:HSJ1141E20	+	39.00	83.49	1.3e+04	AC013205	Drosophila melanogaster
gb_hg1:AC008195	+	39.00	81.94	1.5e+04	AC005365	Homo sapiens chromosome
gb_pr3:HSJ111266	+	39.00	80.56	1.8e+04	AC010912	Homo sapiens chromosome
gb_hg4:AC011957	+	39.00	79.40	2.1e+04	AC008195	Drosophila melanogaster
gb_hg5:AC011702	+	39.00	78.77	2.3e+04	AC080317	Human DNA sequence
gb_in2:AC005286	+	39.00	76.28	2.4e+04	AC011957	Homo sapiens chromosome
gb_pr2:CN01DSC	+	39.00	76.64	3.0e+04	AC011702	Drosophila melanogaster
gb_pr2:HSZ95098	+	38.00	147.71	3.33	AC0121767	Human chromosome 14

gb_in1:PFU6381 + 38.00 134.56 17.95 230 ! AJ006381 Platythreia punctata
gb_pr2:AF130141 - 38.00 126.81 48.54 541 ! AF130141 Fusarium solani f.
gb_bal:UEU131275 - 38.00 122.55 83.80 865 ! AJ131275 uncultured euryarch

seq_name: gb_hg7:AC013772

seq_documentation_block:

LOCUS AC013772 142796 bp DNA HTG 09-DEC-1999
DEFINITION Homo sapiens clone RP11-7024, WORKING DRAFT SEQUENCE, 6 unordered pieces.
ACCESSION AC013772
VERSION AC013772.2 GI:6553994
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE human.
ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Homidae; Homo.

1 (bases 1 to 142796)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens, clone RP11-7024

Unpublished

2 (bases 1 to 142796)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M., Baldwin,J., Barna,N., Beckerly,R., Boguslavsky,L., Boukhalter,B., Brown,A., Castle,A., Colangelo,M., Collins,S., Collamore,A., Cooke,P., Dearellano,K., Dewar,K., Domino,M., Donelan,L., Doyle,M., Ferreira,P., FitzHugh,W., Forrest,C., Funke,R., Gage,D., Galagan,J., Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L., Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J., Lechocsky,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N., McEwan,P., McGuck,A., McKernan,K., McLaughlin,J., Meldrum,J., Morrow,J., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P., Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P., Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X., Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.

Direct Submission

Submitted (15-NOV-1999) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Dec 10, 1999 this sequence version replaced g1:6425750.
All repeats were identified using RepeatMasker:
Smit, A.F.P. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

TITLE

JOURNAL

COMMENT

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WTHR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L2959
Center clone name: 7_O_24

----- Summary Statistics

Sequencing vector: M13; M7815; 100% of reads

Chemistry: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 128649 bases at least Q40

Consensus quality: 137112 bases at least Q30

Consensus quality: 140758 bases at least Q20

Insert size: 142796; sum-of-contigs

Quality coverage: 5.7 in Q20 bases; agarose-fp

Quality coverage: 5.5 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently

* consists of 6 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1

* 2142: contig of 2142 bp in length

* gap of unknown length

*

```

*      2143      4834: contig of 2692 bp in length
*      gap of unknown length
*      4835      9211: contig of 4377 bp in length
*      gap of unknown length
*      9212      25157: contig of 15946 bp in length
*      gap of unknown length
*      25158      66208: contig of 41051 bp in length
*      gap of unknown length
*      66209      142796: contig of 76588 bp in length.

```

FEATURES

```

Location/Qualifiers
1..142796
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-7024"
/clone_lib="RPC1-11 Human Male BAC"
BASE COUNT 47989 a 24580 c 24540 g 45686 t 1 others
ORIGIN

```

```

alignment_scores:
Quality: 47.00      Length: 12
Ratio: 4.273      Gaps: 0
Percent Similarity: 91.667      Percent Identity: 75.000

```

alignment_block:

```

US-08-653-294-18 x AC013772/rev ..

```

```

Align seg 1/1 to reverse of: AC013772 from: 1 to: 142796

```

```

1 TyArgLeuAlaIleArgArgIleLeuLeuArgTvr 12
|||||
13521 TATAGACAGACAGTACAGACAGATTACTATATAT 13486

```

```

seq_name: gb_hgt2:HSBJ655C5

```

seq_documentation_block:

```

LOCUS HSDJ655C5 306999 bp DNA HTG 26-NOV-1999
DEFINITION Homo sapiens chromosome 6 clone RP4-655C5, *** SEQUENCING IN
PROGRESS ***, in unordered pieces.

```

```

ACCESSION AL121956
VERSION AL121956.2 GI:6469398

```

```

KEYWORDS HTG; HTGS_PHASE1.
SOURCE human.

```

ORGANISM

```

Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 306999)

```

```

Sims.S.
Direct Submission

```

REFERENCE

```

AUTHORS Submitted (26-NOV-1999) Sanger Centre, Hinxton, Cambridgeshire,
TITLE CB10 ISA, UK. E-mail enquiries: humquery@sanger.ac.uk clone
JOURNAL requests: clonerequests@sanger.ac.uk

```

COMMENT

```

On Nov 27, 1999 this sequence version replaced gi:6066017.
IMPORTANT: This sequence is unfinished and does not necessarily
represent the correct sequence. Work on the sequence is in
progress and the release of this data is based on the understanding
that the sequence may change as work continues. The sequence may
be contaminated with foreign sequence from E.coli, yeast, vector,
phage etc. Order of segments is not known; 800 n's separate
segments. Contig_ID: 00008 Length: 1036bp

```

```

Contig_ID: 00019 Length: 9963bp
Contig_ID: 00031 Length: 1518bp
Contig_ID: 00039 Length: 1498bp
Contig_ID: 00059 Length: 1840bp
Contig_ID: 00062 Length: 1083bp
Contig_ID: 00088 Length: 1408bp
Contig_ID: 00092 Length: 1477bp
Contig_ID: 00112 Length: 3742bp
Contig_ID: 00142 Length: 1888bp
Contig_ID: 00150 Length: 3371bp
Contig_ID: 00179 Length: 1299bp
Contig_ID: 00183 Length: 3722bp
Contig_ID: 00192 Length: 1260bp

```

```

Contig_ID: 00211 Length: 2606bp
Contig_ID: 00242 Length: 1648bp
Contig_ID: 00248 Length: 4175bp
Contig_ID: 00272 Length: 1781bp
Contig_ID: 00323 Length: 10363bp
Contig_ID: 00355 Length: 1790bp
Contig_ID: 00387 Length: 4242bp
Contig_ID: 00394 Length: 6397bp
Contig_ID: 00404 Length: 1731bp
Contig_ID: 00421 Length: 1266bp
Contig_ID: 00476 Length: 1153bp
Contig_ID: 00481 Length: 980bp
Contig_ID: 00528 Length: 2380bp
Contig_ID: 00555 Length: 2876bp
Contig_ID: 00567 Length: 1458bp
Contig_ID: 00603 Length: 2634bp
Contig_ID: 00605 Length: 1049bp
Contig_ID: 00638 Length: 1196bp
Contig_ID: 00679 Length: 3286bp
Contig_ID: 00695 Length: 4879bp
Contig_ID: 00709 Length: 3049bp
Contig_ID: 00722 Length: 5922bp
Contig_ID: 00725 Length: 1901bp
Contig_ID: 00750 Length: 1420bp
Contig_ID: 00812 Length: 1513bp
Contig_ID: 00892 Length: 1333bp
Contig_ID: 00914 Length: 14732bp
Contig_ID: 00940 Length: 1506bp
Contig_ID: 00964 Length: 1687bp
Contig_ID: 00973 Length: 1044bp
Contig_ID: 00979 Length: 1055bp
Contig_ID: 01017 Length: 3425bp
Contig_ID: 01057 Length: 3223bp
Contig_ID: 01085 Length: 1282bp
Contig_ID: 01125 Length: 3177bp
Contig_ID: 01137 Length: 3871bp
Contig_ID: 01155 Length: 1111bp
Contig_ID: 01212 Length: 1153bp
Contig_ID: 01219 Length: 5468bp
Contig_ID: 01220 Length: 1788bp
Contig_ID: 01238 Length: 3851bp
Contig_ID: 01243 Length: 4555bp
Contig_ID: 01246 Length: 1021bp
Contig_ID: 01256 Length: 1097bp
Contig_ID: 01288 Length: 3286bp
Contig_ID: 01374 Length: 5117bp
Contig_ID: 01396 Length: 2253bp
Contig_ID: 01370 Length: 1884bp
Contig_ID: 01371 Length: 1208bp
Contig_ID: 01374 Length: 6578bp
Contig_ID: 01413 Length: 1372bp
Contig_ID: 01427 Length: 1097bp
Contig_ID: 01430 Length: 2927bp
Contig_ID: 01435 Length: 1190bp
Contig_ID: 01453 Length: 3272bp
Contig_ID: 01487 Length: 1704bp
Contig_ID: 01519 Length: 1071bp
Contig_ID: 01524 Length: 1081bp
Contig_ID: 01530 Length: 2110bp
Contig_ID: 01546 Length: 1126bp
Contig_ID: 01592 Length: 1168bp
Contig_ID: 01630 Length: 8584bp
Contig_ID: 01632 Length: 1827bp
Contig_ID: 01717 Length: 2693bp
Contig_ID: 01740 Length: 1334bp
Contig_ID: 01745 Length: 1861bp
Contig_ID: 01746 Length: 1912bp
Contig_ID: 01776 Length: 2890bp
Contig_ID: 01822 Length: 2552bp
Contig_ID: 01830 Length: 1669bp
Contig_ID: 01853 Length: 1222bp
Contig_ID: 01904 Length: 1994bp
Contig_ID: 01918 Length: 1102bp

```

* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

```

----- Summary Statistics -----
Sequencing vector: M13; M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 116065 bases at least Q40
Consensus quality: 135485 bases at least Q30
Consensus quality: 148288 bases at least Q20
Insert size: 164000; agarose-fp
Insert size: 156243; sum-of-contents
Quality coverage: 3.7 in Q20 bases; agarose-fp
Quality coverage: 3.9 in Q20 bases; sum-of-contents

```

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

```

* 1 4018: contig of 4018 bp in length
* be preserved.

4019	18534:	contig of 14516 bp in	gap of unknown length
*			
*			
*			
*	18535	29198:	contig of 10664 bp in
			gap of unknown length

★ 29199 44793: contig of 15595 bp in length
gap of unknown length

★	44794	65358:	contig of 20565 bp in length
★			gap of unknown length
★	65359	103069:	contig of 37711 bp in length
★			gap of unknown length

* 103070 156243: contig of 53174 bp in length.
Location/Qualifiers
1. .156243
/organism="Homo sapiens"

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-2A8"
/clone_lib="RPCI-11 Human Male BAC"
45948 a 32079c 31296 g 46894 t 26 others

```

[illegible]

Accession	Gene	Size (bp)	Library
U01115	HTG	37459	DNA
U01116	HTG	37459	DNA
U01117	HTG	37459	DNA
U01118	HTG	37459	DNA
U01119	HTG	37459	DNA
U01120	HTG	37459	DNA
U01121	HTG	37459	DNA
U01122	HTG	37459	DNA
U01123	HTG	37459	DNA
U01124	HTG	37459	DNA
U01125	HTG	37459	DNA
U01126	HTG	37459	DNA
U01127	HTG	37459	DNA
U01128	HTG	37459	DNA
U01129	HTG	37459	DNA
U01130	HTG	37459	DNA
U01131	HTG	37459	DNA
U01132	HTG	37459	DNA
U01133	HTG	37459	DNA
U01134	HTG	37459	DNA
U01135	HTG	37459	DNA
U01136	HTG	37459	DNA
U01137	HTG	37459	DNA
U01138	HTG	37459	DNA
U01139	HTG	37459	DNA
U01140	HTG	37459	DNA
U01141	HTG	37459	DNA
U01142	HTG	37459	DNA
U01143	HTG	37459	DNA
U01144	HTG	37459	DNA
U01145	HTG	37459	DNA
U01146	HTG	37459	DNA
U01147	HTG	37459	DNA
U01148	HTG	37459	DNA
U01149	HTG	37459	DNA
U01150	HTG	37459	DNA
U01151	HTG	37459	DNA
U01152	HTG	37459	DNA
U01153	HTG	37459	DNA
U01154	HTG	37459	DNA
U01155	HTG	37459	DNA
U01156	HTG	37459	DNA
U01157	HTG	37459	DNA
U01158	HTG	37459	DNA
U01159	HTG	37459	DNA
U01160	HTG	37459	DNA
U01161	HTG	37459	DNA
U01162	HTG	37459	DNA
U01163	HTG	37459	DNA
U01164	HTG	37459	DNA
U01165	HTG	37459	DNA
U01166	HTG	37459	DNA
U01167	HTG	37459	DNA
U01168	HTG	37459	DNA
U01169	HTG	37459	DNA
U01170	HTG	37459	DNA
U01171	HTG	37459	DNA
U01172	HTG	37459	DNA
U01173	HTG	37459	DNA
U01174	HTG	37459	DNA
U01175	HTG	37459	DNA
U01176	HTG	37459	DNA
U01177	HTG	37459	DNA
U01178	HTG	37459	DNA
U01179	HTG	37459	DNA
U01180	HTG	37459	DNA
U01181	HTG	37459	DNA
U01182	HTG	37459	DNA
U01183	HTG	37459	DNA
U01184	HTG	37459	DNA
U01185	HTG	37459	DNA
U01186	HTG	37459	DNA
U01187	HTG	37459	DNA
U01188	HTG	37459	DNA
U01189	HTG	37459	DNA
U01190	HTG	37459	DNA
U01191	HTG	37459	DNA
U01192	HTG	37459	DNA
U01193	HTG	37459	DNA
U01194	HTG	37459	DNA
U01195	HTG	37459	DNA
U01196	HTG	37459	DNA
U01197	HTG	37459	DNA
U01198	HTG	37459	DNA
U01199	HTG	37459	DNA
U01200	HTG	37459	DNA

AC012150.1 GI:6091632

HTG; HTGS_PHASE1.
human.


```

US-08-653-294-18 x DMRNA3
Align seg 1/1 to: DMRNA3 from: 1 to: 1050

1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||:|||||:
922 TACAGATGTCGATGACGACGATTGCTTACGTTAT 957

seq_name: gb_pr2:MMHLAFHOM
seq_documentation_block:
LOCUS MMHLAFHOM 1056 bp mRNA
DEFINITION M.mullatta HLA-F like mRNA.
ACCESSION Z21819
KEYWORDS HLA-F gene.
SOURCE rhesus monkey.
ORGANISM Macaca mulatta
REFERENCE 1 (bases 1 to 1056)
AUTHORS Bontrop,R.E.
TITLE Characterization of the rhesus macaque (Macaca mulatta) equivalent
of HLA-F
JOURNAL Immunogenetics (1993) In press
REFERENCE 2 (bases 1 to 1056)
AUTHORS Bontrop,R.E.
TITLE Direct Submission
JOURNAL Submitted (16-FEB-1993) Ronald R.E. Bontrop Ph.D, chronic and
infectious diseases, ITRI-TNO, Lange Kleiweg 151, Rijswijk, 2280
HV, The Netherlands
FEATURES
source
1..1056
Location/Qualifiers
/organism="Macaca mulatta"
/isolate="ikm"
/db_xref="taxon:9544"
<1..>1056
10..1056
/codon_start=1
/product="HLA-F like protein"
/protein_id="CAA79885.1"
/db_xref="GI:38569"
/db_xref="SWISS-PROT:P33617"
/translacion="MAPRTLLVLSGALALTETWAGSHSLRYFSTAVSRPGRGSPQVLR
YIAVSYDDTQFLRFDSAAIPRMEPRAPVVEQGPQYWERITGYAKANARTDRVALR
KLLRLYNQSEAGSHTLQMGNGCDMPDGRLLRGYHQAYDKDYISLNEDLRSWTAD
TVARITGRFEAEYAEFRYILEGCELELLRLYENGKELQKLRADPFKAHLHPVS
DREATLRCWALGFYPPDITLTWQRDEEQTDTELVTETRPAGDGTFOKAAVVPVSGE
EQRYTCHVQHEGLPQDLRLWESSQPTIPVIGVAGLAVLVAVVTVGAVVAANVMRRK
SSDRNRGSYQAAM"
mat_peptide
BASE COUNT 214 a 318 c 349 g 175 t
ORIGIN
..
alignment_scores:
Quality: 41.00 Length: 11
Ratio: 3.727 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 63.636
alignment_block:
US-08-653-294-18 x MMHLAFHOM
Align seg 1/1 to: MMHLAFHOM from: 1 to: 1056

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||:|||||:
301 CGAGTGGCCCTGAGGAAGCTGCTCCTCCGCTAC 333

seq_name: gb_htg7:AC017966
seq_documentation_block:
LOCUS AC017966 8849 bp DNA
DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in Ordered
pieces.
ACCESSION AC017966
VERSION AC017966.1 GI:5553224
KEYWORDS HTG; HTGS_PHASE2.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 8849)
AUTHORS Adams,M. and Venter,J.C.
TITLE Direct Submission
JOURNAL Submitted (09-DEC-1999) Celera Genomics, 45 West Gude Drive,
Rockville, MD, USA
COMMENT This sequence was identified as CDM:10212817 by the submitter.
For more information on this record e-mail to fly@celera.com.
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
FEATURES
source
1..8849
Location/Qualifiers
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
BASE COUNT 2604 a 1735 c 1884 g 2626 t
ORIGIN
..
alignment_scores:
Quality: 41.00 Length: 12
Ratio: 3.727 Gaps: 0
Percent Similarity: 91.667 Percent Identity: 58.333
alignment_block:
US-08-653-294-18 x AC017966
Align seg 1/1 to: AC017966 from: 1 to: 8849

1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||:|||||:
844 TACAGATGTCGATGACGACGATTGCTTACGTTAT 879

seq_name: gb_htg5:AC013927
seq_documentation_block:
LOCUS AC013927 12771 bp DNA
DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in Ordered
pieces.
ACCESSION AC013927
VERSION AC013927.1 GI:5437408
KEYWORDS HTG; HTGS_PHASE2.
SOURCE fruit fly.
ORGANISM Drosophila melanogaster
Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE 1 (bases 1 to 12771)
AUTHORS Adams,M. and Venter,J.C.
TITLE Direct Submission
JOURNAL Submitted (16-NOV-1999) Celera Genomics, 45 West Gude Drive,
Rockville, MD, USA
COMMENT This sequence was identified as CDM:10213548 by the submitter.
For further information on this sequence e-mail to fly@celera.com.
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
FEATURES
source
1..12771
Location/Qualifiers
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"

```

BASE COUNT 3823 a 2428 c 2497 g 4023 t
ORIGIN

alignment_scores:
Quality: 41.00 Length: 11
Ratio: 4.100 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 81.818

alignment_block:

US-08-653-294-18 x AC013927/rev ..

Align seg 1/1 to reverse of: AC013927 from: 1 to: 12771

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
2697 CGGTGGCAATCATCGCAATTTATTGCAATAT 2665

seq_name: gb_bai:MLCB2533

seq_documentation_block: MLCB2533 40245 bp DNA BCT 27-AUG-1999

LOCUS MLCB2533 40245 bp DNA BCT 27-AUG-1999
DEFINITION Mycobacterium leprae cosmid B2533.
ACCESSION AL035310
VERSION AL035310.1 GI:4200258

KEYWORDS
ansp: ATP-dependent RNA helicase; ATP-phosphoribosyl transferase;
ATPase; hisG 5-methyltetrahydrofolate-homocysteine methyl
transferase; hisI; L-asparagine permease; meth; mtb;
phosphoribosyl-AMP cyclohydrolase; prcA; prcB; proteasome alpha
subunit; proteasome beta subunit; protein translocation system;
pseudogene; RLEP; sec-independent.

SOURCE

Mycobacterium leprae.
Mycobacterium leprae
Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
Actinomycetales; Corynebacterineae; Mycobacteriaceae;
Mycobacterium.

REFERENCE
1 (bases 1 to 40245)
Eiglmeier, K., Honore, N., Woods, S.A., Caudron, B. and Cole, S.T.
Use of an ordered cosmid library to deduce the genomic organization
of Mycobacterium leprae
Mol. Microbiol. 7 (2), 197-206 (1993)

JOURNAL

MEDLINE

REFERENCE

2 (bases 1 to 40245)

Hamlin, N. and Churcher, C.M.

Unpublished

3 (bases 1 to 40245)

James, K.D., Parkhill, J., Barrell, B.G. and Rajandream, M.A.

Direct Submission

Submitted (26-JAN-1998) Mycobacterium leprae sequencing project,

Sanger Centre, Wellcome Trust Genome Campus, Hinxton, Cambridge

CB10 LSA E-mail: barrells@sanger.ac.uk Cosmids supplied by Dr.

Stewart T. Cole, [3] Unite de Genetique Moleculaire Bacterienne,

Institut Pasteur, 28 rue du Docteur Roux, 75724 Paris Cedex 15,

France Requests for cosmids should be sent to Karin Eiglmeier

(keig@pasteur.fr)

Notes:

The Sanger Centre is funded to complete the sequence of M. leprae

by the Heiser Program for Research in Leprosy and Tuberculosis of

The New York Community Trust.

Work in Paris is supported by the Heiser Trust, the Association

Francaise Raoul Follereau and the Groupement de Recherches et des

Etudes des Genomes (GIP-GREG).

Details of M. leprae sequencing at the Sanger Centre are available

on the World Wide Web.

(URL, <http://www.sanger.ac.uk/Projects/>)

CDS are numbered using the following system eg MLCB33.01c. ML (M.

leprae), CB33 (cosmid name), .01 (first CDS), c (complementary

strand).

The more significant matches with motifs in the PROSITE database

are also included but some of these may be fortuitous. The length

in codons is given for each CDS.

Usually the highest scoring match found by fasta -o is given for

CDS which show significant similarity to other CDS in the database.

The position of possible ribosome binding site sequences are given where these have been used to deduce the initiation codon. All CDS over 100 codons have been analysed. Gene prediction is based on positional base preference in codons especially where there is an increase in the observed/expected third position G + C. CAUTION: We may not have predicted the correct initiation codon. Where possible we choose an initiation codon (atg, gtg, or ttg) which is preceded by an upstream ribosome binding site sequence (optimally 5-13bp before the initiation codon). If this cannot be identified we choose the most upstream initiation codon.

FEATURES

Source

1. .40245

/organism="Mycobacterium leprae"

/db_xref="taxon:1769"

/clone="cosmid B2533"

complement(1. .1099)

/gene="ansp"

complement(1. .31687)

/note="overlap with EMBL:ML017 cosmid B2126 from 1 to

31682. There are 16 conflicts between this sequence and

ours. In each case our sequence has been checked and is

thought to be correct"

complement(<1. .1099)

/gene="ansp"

/note="MLCB2533.01c, ansp, probable L-asparagine permease,

partial CDS, len: >366 aa; highly similar to many

amino-acid permeases e.g. ANSP_SALTY (EMBL:U04851)

S.typhimurium Ansp, L-asparagine permease (L-asparagine

transport protein) (497 aa) fasta scores: opt: 1508

z-score: 1696.8 E(-): 0, 61.2% identity in 366 aa overlap.

Equivalent to M.tuberculosis RV2127, ansp (MTCY13E10.26,

85.7% identity in 356 aa overlap). Also similar to

75.1% identity in 345 aa overlap). Probable integral

membrane protein, contains PS00218 Amino acid permease

signature. Pfam match to entry PF00324 aa_permeases, amino

acid permease. Annotated as ORF TR:Q49801, designated

arop2 in M.leprae cosmid EMBL:U00017"

/codon_start=1

/transl_table=11

/product="putative L-asparagine permease"

/protein_id="CAA22915.1"

/db_xref="GI:4200259"

/translation="MATLAESPESKSGASRAGVLGEAGYHKGKLPQLOMIGIGAI

GTGLFLGAGRLAKAGPLFLVAVGVFLILRALGELVLRHSSGSFVSAREFF

GKAAVVGWLYFLDQAMTAIVDTATATYLRHTIFTALPQNTLALLAVLVNML

ISVQFGELEFWAALKYCALMAFLVGTIFLGGYPVDGHTGLSLMTSHGLEPTG

VAQLVSSGVNFAYANVELVGTAGETVEPKKIMPRAINSIAIAFYVGSVILLA

LLLPFSAPKASESPVTFKSGVFGAGDLNIVLTALSSLSNAGLIATGRVMSIA

INGSGPKFTARMSKNGVPYGGILLAAVICLG"

1. .1168

/note="1168 bp perfect direct repeat"

complement(2. .991)

/gene="ansp"

/note="Pfam match to entry PF00324 aa_permeases, Amino

acid permease, score 245.80, E-value 6e-70"

438

/gene="ansp"

/note="ambiguous base T /G"

complement(833. .925)

/gene="ansp"

/note="PS00218 Amino acid permease signature"

1223

/note="conflict: T is TG EMBL:ML017"

complement(1249. .2766)

/gene="ansp2"

complement(1249. .2766)

/gene="ansp2"

/note="MLCB2533.02c, ansp2, probable L-asparagine

```

permease, len: 505 aa; highly similar to many amino-acid
permeases e.g. ANSP_SALTY (EMBL:U04851) S.typhimurium
Ansp, L-asparagine permease (L-asparagine transport
protein) (497 aa), fasta scores:opt: 1891 z-score: 2218.9
E(): 0, 58.9% identity in 477 aa overlap. Equivalent to
M.tuberculosis Rv2127 (MTCY261.26, 83.7% identity in 485
aa overlap). Also similar to M.tuberculosis permease
Rv0346c (MTCV1310.06c, 69.8% identity in 473 aa overlap).
Probable integral membrane protein, contains PS00218 Amino
acid permeases signature. Pfam match to entry PF00324
aa_permeases, amino acid permease. Annotated as ORF
TR:Q49802, designated lysp in M.leprae cosmid EMBL:U00017"
/codon_start=1
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/protein_id="CAA22916.1"
/db_xref="GI:4200260"
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GTGLFGAGRLAKAGFLFYAVCGVFLILRALGELVLRHPSGGSFVSAREFF
GEKAYVGLWFLDAMTAIVDTTATYTLRWITFTALPQWTLALLAVLYVMNL
ISVMEFGELEFNAALIKVCAALFLVVGTFILGGRYPVDGHTGLSWTSHGGLFPTG
VAPLIVSSGVMPAYAAVELVGTAGETVEPKIMPRAINSVIARTAIYVGSVILLA
LLPYSAFASPSPFYFSGYFCAGDLNIVLTALSSNLAGLYATGRVMSIA
INGSGPFTARSKNGVPYGGILLAAVICLGVNALNFPQGAFEIVLSVAGHIIAG
WGTIVLCQLRLHKMAGIMRFRMRPLAPYSGYTLTFLFVLYVMAFDKPIGTWT
VASLIVIPALTAGWTSIRKRVNTIARERMGYTGPPAIANPVPQPSRSHSNP"
complement(1354..2691)
/genes="ansp2"
/notes="Pfam match to entry PF00324 aa_permeases, Amino
acid permease, score 501.80, E-value 5.1e-147"
1668..2835
/notes="1168 bp perfect direct repeat"
2104
/genes="ansp2"
/notes="conflict: C is CT in EMBL:ML017"
complement(2500..2592)
/genes="ansp2"
/notes="PS00218 Amino acid permeases signature"
complement(2903..3856)
/genes="MLCB2533.03c"
complement(2903..3856)
/genes="MLCB2533.03c"
/notes="MLCB2533.03c"
similar to M.tuberculosis hypothetical protein Rv2125
(MTCY261.21) (EMBL:Z97559) (292 aa), fasta scores: opt:
1648 z-score: 2382.2 E(): 0, 84.1% identity in 290 aa
overlap. Also some similarity to M.leprae hypothetical
protein TR:Q49847 (297% identity in 279 aa overlap).
Annotated as ORF TR:Q49797, hypothetical protein in
M.leprae cosmid EMBL:U00017"
/codon_start=1
/transl_table=11
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/protein_id="CAA22917.1"
/db_xref="GI:4200261"
/db_xref="SPTREMBL:Q49797"
/translation="MPPHRAVTRHASSALKPYADSVTLRDGDPDRGALPELHNTVVV
AAFEQNDASDAASGALEHLNAWEADPIVEIDDEAYDYQVNRPIRVQDVGTVREL
WPMRLISYCRPGSDRNVLMHGVEPNRWRFTCTELLTIADRLNDVTIVILGALLAD
THPRVPVSGAAYSARRFEGLEETRYEGPTGIAGVFDACVAARIAPVFWAAYV
HYVSHPNPKATVALLRVEDVLDVEVLADLPQAEDEWEQAITAEADDEALAEYVH
SLEQRDAEVDVNDALGKIDGDALEEFYLRRLRRPGFGR"
4004..7555
/genes="meth"
4004..7555
/genes="meth"
/notes="MLCB2533.04, meth, probable
5-methyltetrahydrofolate-homocysteine methyltransferase,
len: 1183 aa; similar to many members of vitamin-B12
dependent methionine synthase family e.g. METH_ECOLI
(EMBL:X16584) E.coli meth (1226 aa), fasta scores: opt:
1617 z-score: 1000.7 E(): 0, 31.6% identity in 1228 aa
overlap. Equivalent to M.tuberculosis Rv2124c

```

```

(MTCY261.20c, 88.7% identity in 1183 aa overlap).
Annotated as METH_MYCLE, designated meth2 in M.leprae

alignment_scores:
  Quality: 41.00      Length: 12
  Ratio: 4.556       Gaps: 0
  Percent Similarity: 75.000  Percent Identity: 75.000

alignment_block:
US-08-653-294-18 x MLCB2533 ..
Align seg 1/1 to: MLCB2533 from: 1 to: 40245
1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
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17534 TATCGATTGGCAATCGCAGATATCTGCACGGTAT 17569

seg_name: gb_bai:U00017

seg_documentation_block:
LOCUS      U00017      42157 bp      DNA      BCT      01-MAR-1994
DEFINITION  Mycobacterium leprae cosmid B2126.
ACCESSION  U00017
VERSION    U00017.1  GI:466994
KEYWORDS
SOURCE     Mycobacterium leprae.
ORGANISM   Mycobacterium leprae
Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
Actinomycetales; Corynebacterineae; Mycobacteriaceae;
Mycobacterium.
REFERENCE  1 (bases 1 to 42157)
AUTHORS   Smith,D.R.
JOURNAL   Unpublished
REFERENCE  2 (bases 1 to 42157)
AUTHORS   Robison,K.
TITLE     Direct Submission
JOURNAL
REFERENCE  3 (bases 1 to 42157)
AUTHORS   Robison,K.
TITLE     Direct Submission
JOURNAL
COMMENT   Submitted (01-MAR-1994) Department of Genetics, Harvard Medical
School, 200 Longwood Avenue, Boston MA 02115
On Mar 31, 1994 this sequence version replaced gi:414223.
This sequence data was produced by the Genome Sequencing Center
located at Collaborative Research Incorporated (1365 Main St.,
Waltham MA, 02159). 617-487-7979. Please contact Doug Smith
(smith@eric.com). The annotation should be considered
preliminary and incomplete.
FEATURES             Location/Qualifiers
     1..42157
     /organism="Mycobacterium leprae"
     /db_xref="taxon:1769"
     complement(133..1086)
     /note="match to yigU and yigV E.coli; B2126_C1_183"
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LIIISLAAVTTTIFGFIWHSIFGLSIGELWRPVCYSLPOSARADISPDQCRLLA
TAPDQFMRLTKVGMAGVILASPVWFQQLWAFITPGLYTKERRFTVAFVPAALFA
GTGVLAYLIVKALGFLIIVSGVQVYALSGDRYFGLNLLVWFGVGFEPPLLIVML
NIAGLTYQRLKSRWRGLIFAMVFAAVTFPGSDPFSMTALGAALTLLLEALQLVRL
HDKRRVKHEALIDAEASVIEPPSPISPERTATRSDDVT"
     complement(1132..1398)
     /note="match to yigT E.coli; B2126_C1_182"
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     /transl_table=11
     /product="u2126b"
     /protein_id="AAAL1790.1"
     /db_xref="GI:467005"

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CDS
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/protein_id="AA17189.1"
/db_xref="GI:467004"

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QVQAPALAYVRNAVAALSYQVTAQ"
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/transl_table=11
/product="B2126_C2_220"
/protein_id="AA17199.1"
/db_xref="GI:467014"

CDS
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NQLWCGLPVGGDLIDFEFGSDTIEVTSAGIDRPLQLTSPEAIGLLVALRALANI
PGVDPPEAVRSIAIKIEAAVVMGNEATGSVSDVTRFSESHAVAAVRAAVRNKQAL
VTDYTSAGDHTLSTRIVDP IRVLLVGDHSYLEAMSREAEGVRLFRDRIYVARELDEP
AAAPTVRRCOTHRSSMTTRCCRRRCG"
complement(2441..3436)
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/transl_table=11
/product="B2126_C3_266"
/protein_id="AA17208.1"
/db_xref="GI:467023"

CDS
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SMFPRDNELRDGLGPILEVGVKVALDSEGRYINRDYALPPVELITDEAAVAVAT
QWESQELTQAGALLKRAAGVDIDPLDTPVJASSGSSVSLRGSDDFLSLLGAI
GSRQVAPYPRSAEPTMRNVPEWITENSCWYLVGHDCDRNATFRLSLGSE
VAPIGPAGAVTVDGRLRIVSDAVSVTGATARVWVVDGRATALRHAGRPAGVR
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complement(3443..3841)
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/db_xref="GI:467022"

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GVFDLLQKGLAARTDEDIADAVNHPPQTRARLRGEFISAQAAGRDTVDWVHL
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/product="B2126_C2_219"
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/db_xref="GI:467013"

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SVLLPFLVTLQICGACKVLOTPKAATFCLQSRAEHIEWGVSSATTSRPIINTRDE
PHADAEKRLRHLVIVGDSNMCETTTMLKVGTAALMLEWVGVPFRDPSLONPRAIR
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/protein_id="AA17206.1"
/db_xref="GI:467021"

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LVFVRPATKMGVLRTMSESDAIGQVADWLTPGSVLMCLTVRHAFLVKILV"
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/transl_table=11
/product="B2126_C2_217"
/protein_id="AA17197.1"
/db_xref="GI:467012"

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AAFEDEELVRLSCGHERRSFLSHRLAPHLPARLITIHENRRERRAAVFR"
complement(7253..8050)
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NPSRLQKISELYDRVGVFAAAGKFNEDFNLRGGIQPADRGYAYDRDDYTGQLANV
YAQTLGTTFTEQAKPYEVLCAVEAVHYGHTKPELYRITYIDGSIINDEPHFMVGGT
ESTANALKEYSANASLTDLGIAVAARLAGSADAAGSDPTLGVASLEVAVLDANRP
RRAPRIIGSGLEALLREKDSKSGKGAQNPKGARDSKNSKSYGESTD"
complement(8047..8922)
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AGDIHAPDQSGAGRIYSDAAGWNIEEGYQSGSGSIFAKSSIKKLYSOVSDADS
ALRVAIEALYDAADDDSATGGDLVRGIYPTAVTIGAEGAAEVTESRIAREIIES
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APETCDLDAVIMDKAGERWEAAARHVASVPGAAKQLKNNVDGKAGSYGAHENTL
MSRQTPSAIAGTLPELVSRQVVTGSGRVIGPAGDPGFQLSQSDYIEVEVGLT
TLKGGIINTREDEHADRYRLRHVIYGDANLAETSTYKLTGLTLDLIEGPHVG
IDLTLALRPVHVAHAI SRDASRLATVLDGRELGTALQRIYLDRAKLVDSRDP
DRAADVVTWVHVDOLERDMDCAELLDPAKRLLEGFRORENLNWAPRLHVD
LQYSDVRLKGLYNLVARSGMKELVNEHQVLRVNNPPTDTRAYFEGECLRRFSADI
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/protein_id="AA17203.1"
/db_xref="GI:467018"

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CDS          14353..14763

alignment_scores:
  Quality: 41.00      Length: 12
  Ratio: 4.556       Gaps: 0
  Percent Similarity: 75.000  Percent Identity: 75.000

alignment_block:
US-08-653-294-18 x U00017/rev ...

Align seg 1/1 to reverse of: U00017 from: 1 to: 42157

      1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
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14153 TATCGATTGGCAAGTCGCAGATATCTGCACGGTAT 14118

seq_name: gb_htg6:AC007807

seq_documentation_block:
LOCUS      AC007807 130536 bp DNA HTG 23-NOV-1999
DEFINITION Drosophila melanogaster chromosome 3 clone BACR01E04 (D714) RPCI-98
            01.E.4 map 89E-89E strain y; cn bw sp, *** SEQUENCING IN PROGRESS
            ***, 99 unordered pieces.
ACCESSION  AC007807
VERSION     AC007807.4 GI:6466499
KEYWORDS   HTG: HTGS_PHASE1.
SOURCE     fruit fly.
ORGANISM   Drosophila melanogaster
            Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
            Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
            Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
REFERENCE  1 (bases 1 to 130536)
AUTHORS   Celniker,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
            Butenhoff,C., Champe,M., Chavez,C., Chew,M., Ciesiolka,L.,
            Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
            Hinkle,A., Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K.,
            Kearney,L., Lee,B., Lewis,S., Li,P., Ling,H., Moshrefi,A.R.,
            Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S., Pfeiffer,B.,
            Richards,S., Sethi,H., Svirskas,R.R., Wan,K.H., Webster,D.,
            Woolley,P., Yang,S., Yee,M., Yu,C. and Rubin,G.M.
            Sequencing of Drosophila melanogaster
            Unpublished
            2 (bases 1 to 130536)
AUTHORS   Celniker,S.E., Agbayani,A., Arcaina,T.T., Baxter,E., Blazej,R.G.,
            Butenhoff,C., Champe,M., Chavez,C., Chew,M., Ciesiolka,L.,
            Doyle,C.M., Farfan,D.E., Galle,R., George,R.A., Harris,N.L.,
            Hoskins,R.A., Houston,K.A., Hummasti,S.R., Karra,K., Kearney,L.,
            Kim,E., Lee,B., Lewis,S., Li,P., Lomotan,M.A., Mazda,P.,
            Moshrefi,A.R., Moshrefi,M., Nixon,K., Pacleb,J.M., Park,S.,
            Pfeiffer,B., Poon,L., Sequeira,A., Sethi,H., Snir,E.,
            Svirskas,R.R., Wan,K.H., Weinburg,T., Zhang,R., Zieran,L.L. and
            Rubin,G.M.
            Direct Submission
            Submitted (14-JUN-1999) Drosophila Genome Center, Lawrence Berkeley
            Laboratory, MS 64-121, Berkeley, CA 94720, USA
COMMENT    On Nov 23, 1999 this sequence version replaced gi:5670524.
            For further information about this sequence, including its location
            and relationship to other sequences, please visit our sequence
            archive web site (http://www.fruitfly.org/sequences/) or send email
            to bdp@fruitfly.berkeley.edu. All contigs in this submission meet
            the following cutoffs: length >= 200 bases.
            * NOTE: This is a 'working draft' sequence. It currently
            * consists of 99 contigs. The true order of the pieces
            * is not known and their order in this sequence record is
            * arbitrary. Gaps between the contigs are represented as
            * runs of N, but the exact sizes of the gaps are unknown.
            * This record will be updated with the finished sequence
            * as soon as it is available and the accession number will
            * be preserved.
            * 1 524: contig of 524 bp in length
            * 525 604: gap of unknown length
            * 605 1174: contig of 570 bp in length
            * 1175 1254: gap of unknown length
            *
            1255 1717: contig of 463 bp in length
            * 1718 gap of unknown length
            * 1798 2374: contig of 577 bp in length
            * 2375 gap of unknown length
            * 2455 3028: contig of 572 bp in length
            * 3027 gap of unknown length
            * 3107 3704: contig of 598 bp in length
            * 3705 gap of unknown length
            * 3785 4384: contig of 579 bp in length
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            * 4444 4935: contig of 492 bp in length
            * 4936 gap of unknown length
            * 5016 5662: contig of 646 bp in length
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            * 5742 6375: contig of 633 bp in length
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            * 6455 7007: contig of 553 bp in length
            * 7008 gap of unknown length
            * 7088 7728: contig of 641 bp in length
            * 7729 gap of unknown length
            * 7809 8383: contig of 574 bp in length
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            * 9115 9914: contig of 800 bp in length
            * 9915 gap of unknown length
            * 9995 11174: contig of 1180 bp in length
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            * 11255 12095: contig of 840 bp in length
            * 12095 gap of unknown length
            * 12175 12955: contig of 780 bp in length
            * 12955 gap of unknown length
            * 13034 13830: contig of 796 bp in length
            * 13831 gap of unknown length
            * 13910 15091: contig of 1181 bp in length
            * 15091 gap of unknown length
            * 15172 15977: contig of 806 bp in length
            * 15978 gap of unknown length
            * 16057 16966: contig of 909 bp in length
            * 16967 gap of unknown length
            * 17046 18238: contig of 1192 bp in length
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            * 18319 18875: contig of 557 bp in length
            * 18876 gap of unknown length
            * 18956 19682: contig of 727 bp in length
            * 19683 gap of unknown length
            * 19763 20523: contig of 761 bp in length
            * 20524 gap of unknown length
            * 20604 21293: contig of 690 bp in length
            * 21294 gap of unknown length
            * 21374 22036: contig of 663 bp in length
            * 22037 gap of unknown length
            * 22117 22908: contig of 790 bp in length
            * 22909 gap of unknown length
            * 22987 23813: contig of 827 bp in length
            * 23814 gap of unknown length
            * 23894 25073: contig of 1179 bp in length
            * 25073 gap of unknown length
            * 25133 26366: contig of 1213 bp in length
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            * 26446 27655: contig of 1209 bp in length
            * 27655 gap of unknown length
            * 27735 29379: contig of 1645 bp in length
            * 29380 gap of unknown length
            * 29459 30445: contig of 986 bp in length
            * 30446 gap of unknown length
            * 30526 31238: contig of 713 bp in length
            * 31239 gap of unknown length
            * 31319 32321: contig of 1003 bp in length
            * 32322 gap of unknown length
            * 32401 33223: contig of 822 bp in length
            * 33224 gap of unknown length
            * 33303 34356: contig of 1053 bp in length
            * 34356 gap of unknown length

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* 34357 34436: gap of unknown length
* 34437 contig of 1150 bp in length
* 35586: contig of 1150 bp in length
* 35587 gap of unknown length
* 35667 36738: contig of 1072 bp in length
* 36739 gap of unknown length
* 36818: contig of 791 bp in length
* 37609: contig of 791 bp in length
* 37610 gap of unknown length
* 37689: gap of unknown length
* 37690 contig of 671 bp in length
* 38360: contig of 671 bp in length
* 38361 gap of unknown length
* 38440: gap of unknown length
* 39809: contig of 1369 bp in length
* 39810 gap of unknown length
* 39889: gap of unknown length
* 41023: contig of 1134 bp in length
* 41024 gap of unknown length
* 41103: gap of unknown length
* 41104 contig of 1874 bp in length
* 42977: contig of 1874 bp in length
* 43057: gap of unknown length
* 43058 contig of 1729 bp in length
* 44786: contig of 1729 bp in length
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* 44866: gap of unknown length
* 46003: contig of 1137 bp in length
* 46003: gap of unknown length
* 46083: gap of unknown length
* 46084 contig of 1208 bp in length
* 47291: contig of 1208 bp in length
* 47371: gap of unknown length
* 47372 contig of 1128 bp in length
* 48459: contig of 1128 bp in length
* 48500 gap of unknown length
* 48580 contig of 1861 bp in length
* 50440: gap of unknown length
* 50520: gap of unknown length
* 50521 contig of 2327 bp in length
* 52848: gap of unknown length
* 52848 contig of 1731 bp in length
* 52828 gap of unknown length
* 54738: gap of unknown length
* 54739 contig of 1904 bp in length
* 56642: gap of unknown length
* 56643 gap of unknown length
* 56723 contig of 1189 bp in length
* 57911: contig of 1189 bp in length
* 57921 gap of unknown length
* 57922 contig of 1253 bp in length
* 59324: gap of unknown length
* 59325 contig of 2054 bp in length
* 61378: gap of unknown length
* 61379 contig of 1956 bp in length
* 61459 gap of unknown length
* 63415 contig of 1179 bp in length
* 63495 gap of unknown length
* 64673: contig of 1179 bp in length
* 64674 gap of unknown length
* 64754 contig of 1370 bp in length
* 66123: gap of unknown length
* 66203: gap of unknown length
* 66204 contig of 1218 bp in length
* 67421: gap of unknown length
* 67501: gap of unknown length
* 67502 contig of 1286 bp in length
* 68787: gap of unknown length
* 68788 contig of 1864 bp in length
* 68868 gap of unknown length
* 70732 contig of 2217 bp in length
* 70812: gap of unknown length
* 73028: contig of 1954 bp in length
* 73029 gap of unknown length
* 73063 contig of 1954 bp in length
* 75142: gap of unknown length
* 75143 contig of 2483 bp in length
* 77626 gap of unknown length
* 77705: gap of unknown length
* 77706 contig of 2174 bp in length
* 79879: gap of unknown length
* 79880 gap of unknown length
* 79881 contig of 1636 bp in length
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* 81596: contig of 1636 bp in length
* 81596 gap of unknown length
* 81676 contig of 1960 bp in length
* 83636: gap of unknown length
* 83715: gap of unknown length
* 83716 contig of 1736 bp in length
* 85452: gap of unknown length
* 85452 contig of 2474 bp in length
* 85332 gap of unknown length
* 88006: gap of unknown length
* 88006 contig of 1954 bp in length
* 90039: gap of unknown length
* 90040 gap of unknown length
* 90120 contig of 2225 bp in length
* 92345: gap of unknown length
* 92345 contig of 3090 bp in length
* 92425 gap of unknown length
* 95315: contig of 3291 bp in length
* 95315 gap of unknown length
* 98885: contig of 3291 bp in length
* 98886 gap of unknown length

alignment_scores:
Quality: 41.00 Length: 11
Ratio: 4.100 Gaps: 0
Percent similarity: 90.909 Percent identity: 81.818

alignment_block:
US-08-653-294-18 x AC007807 ...

Align seg 1/1 to: AC007807 from: 1 to: 130536

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||||
83832 CGGCTGGCAATCATCGCAATTTATTCATAT 83864

seq_name: gb_hgt4:AC010564

seq_documentation_block:

LOCUS AC010564 143914 bp DNA HTG 16-OCT-1999
DEFINITION Drosophila melanogaster chromosome 3L/62A1 clone RPC198-2701, ***
SEQUENCING IN PROGRESS ***, 89 unordered pieces.

ACCESSION AC010564

VERSION AC010564.4 GI:5917942

KEYWORDS HTG; HTGS_PHASE1.

SOURCE fruit fly.

ORGANISM

Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;

Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;

Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE

AUTHORS

1 (bases 1 to 143914)
Muzny, D.M., Adams, C., Bailey, M., Barbara, J., Blankenburg, K.,
Bodda, B., Bouck, J., Bowie, S., Brooks, A., Buhay, C., Bunac, C.,
Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C.,
David, R., Delgado, O., Deshazo, D., Ding, Y., Domah-Rashid, N.,
Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D.,
Forcum-Tansey, J., Frantz, P., Ganesh, R., Gorrell, J.H., Gorrell, L.L.,
Guevara, W., Harris, K., Hernandez, J., Hodgson, A., Hogues, M.,
Holloway, C., Hosak, H., Jackson, L.E., Jackson, L., Jia, Y., Jones, M.,
Kelly, S., Kondejowski, N., Kong, Y., Kovar, C., Leal, B., Li, Z.,
Lichtarge, O., Liu, J., Liu, W., Logan, O., Lu, J., Lucier, R.,
Martin, R., Martinez, C., McLeod, M.P., Mei, G., Morgan, M., Morris, S.,
Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Oswal, G.,
Parish, B., Paxton, S., Payton, B., Perez, L., Pu, L., Quiles, M.,
Reiter, D., Rives, M., Samuel, S., Say, J., Scherer, S., Shah, E.,
Shen, H., Simon, M., Sparks, A., Stamps, A., Sucgang, R., Taber, P.,
Taylor, T., Vasquez, L., Vinson, R., Vo, O., Wabwah, M., Watlington, S.,
Weinstock, G., Weinstock, I.R., Williamson, A., Worley, K., Wren, J.,
Wrenford, G., Yu, W., Zhou, X., Nelson, D. and Gibbs, R.

Direct Submission

Unpublished

2 (bases 1 to 143914)

Worley, K.C.

Direct Submission

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

On Sep 22, 1999 this sequence version replaced gi:5916428.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 89 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 831: contig of 831 bp in length
* 832 1645: contig of 814 bp in length
* 1646 2519: contig of 874 bp in length
* 2520 3350: contig of 874 bp in length
* 3351 4206: contig of 856 bp in length
* 4207 5469: contig of 1263 bp in length
* 5470 6313: contig of 844 bp in length
* 6314 7273: contig of 960 bp in length

```
* 7274 8101: contig of 828 bp in length
* 8102 8941: contig of 840 bp in length
* 8942 10144: contig of 1203 bp in length
* 10145 11019: contig of 875 bp in length
* 11020 11825: contig of 806 bp in length
* 11826 13085: contig of 1260 bp in length
* 13086 13937: contig of 852 bp in length
* 13938 14752: contig of 815 bp in length
* 14753 15973: contig of 1220 bp in length
* 15973 16571: contig of 599 bp in length
* 16572 17184: contig of 613 bp in length
* 17185 18018: contig of 834 bp in length
* 18019 18837: contig of 819 bp in length
* 18838 20391: contig of 1554 bp in length
* 20392 21252: contig of 861 bp in length
* 21253 21970: contig of 718 bp in length
* 21971 23026: contig of 1056 bp in length
* 23027 24541: contig of 1515 bp in length
* 24542 25727: contig of 1185 bp in length
* 25728 26570: contig of 844 bp in length
* 26571 27681: contig of 1111 bp in length
* 27682 29230: contig of 1549 bp in length
* 29231 30635: contig of 1405 bp in length
* 30636 31757: contig of 1122 bp in length
* 31758 32899: contig of 1142 bp in length
* 32900 34092: contig of 1193 bp in length
* 34093 34928: contig of 836 bp in length
* 34929 35705: contig of 777 bp in length
* 35706 36891: contig of 1186 bp in length
* 36892 38403: contig of 1512 bp in length
* 38404 39398: contig of 1535 bp in length
* 39399 41157: contig of 1219 bp in length
* 41158 42005: contig of 848 bp in length
* 42006 43465: contig of 1480 bp in length
* 43466 44890: contig of 1423 bp in length
* 44891 45926: contig of 1036 bp in length
* 45927 47498: contig of 1572 bp in length
* 47499 49100: contig of 1602 bp in length
* 49101 50371: contig of 1271 bp in length
* 50372 51976: contig of 1605 bp in length
* 51977 54050: contig of 2074 bp in length
* 54051 56027: contig of 1977 bp in length
* 56028 57411: contig of 1384 bp in length
* 57412 58415: contig of 1404 bp in length
* 58416 60132: contig of 1317 bp in length
* 60133 61360: contig of 1228 bp in length
* 61361 63494: contig of 2134 bp in length
* 63495 64634: contig of 1140 bp in length
* 64635 65855: contig of 1221 bp in length
* 65856 66834: contig of 979 bp in length
* 66835 68045: contig of 1211 bp in length
* 68046 68873: contig of 828 bp in length
* 68874 70019: contig of 1146 bp in length
* 70020 72323: contig of 2304 bp in length
* 72324 73759: contig of 1436 bp in length
* 73760 75834: contig of 2075 bp in length
* 75835 78389: contig of 2555 bp in length
* 78390 80840: contig of 2451 bp in length
* 80841 82111: contig of 1271 bp in length
* 82112 84052: contig of 1941 bp in length
* 84053 86110: contig of 2058 bp in length
* 86111 87488: contig of 1378 bp in length
* 87489 89108: contig of 1620 bp in length
* 89109 91457: contig of 2349 bp in length
* 91458 92907: contig of 1450 bp in length
* 92908 95337: contig of 2430 bp in length
* 95338 97095: contig of 1758 bp in length
* 97096 100091: contig of 2996 bp in length
* 100092 102072: contig of 1981 bp in length
* 102073 104497: contig of 2425 bp in length
* 104498 107508: contig of 3011 bp in length
* 107509 109598: contig of 2090 bp in length
* 109599 112405: contig of 2807 bp in length
```

```
* 112406 115204: contig of 2799 bp in length
* 115205 118317: contig of 3113 bp in length
* 118318 121885: contig of 3568 bp in length
* 121886 125221: contig of 3336 bp in length
* 125222 129338: contig of 4117 bp in length
* 129339 132811: contig of 3473 bp in length
* 132812 137892: contig of 5081 bp in length
* 137893 143914: contig of 6022 bp in length.
```

FEATURES

source

```
1..143914
/organism="Drosophila melanogaster"
/db_xref="taxon:7227"
/chromosome="3L/62A1"
/clone="RPG198-2701"
```

```
BASE COUNT 38720 a 33523 c 32072 g 39042 t 557 others
ORIGIN
```

alignment_scores:

```
Quality: 41.00 Length: 12
Ratio: 3.727 Gaps: 0
Percent Similarity: 91.667 Percent Identity: 58.333
```

alignment_block:

```
US-08-653-294-18 x AC010564 ..
```

```
Align seg 1/1 to: AC010564 from: 1 to: 143914
```

```
1 TyArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
```

```
|||||:|||||:|||||:|||||:|||||:|||||
```

```
58362 TACAGATGTCGATGCAACGATTGCTTACGTAT 58397
```

```
seq_name: gb_htg2:AC005047
```

seq_documentation_block:

```
LOCUS AC005047 159103 bp DNA HTG 12-JUN-1998
DEFINITION Homo sapiens clone RG014E15, *** SEQUENCING IN PROGRESS ***, 2
unordered pieces.
```

```
ACCESSION AC005047
```

```
VERSION AC005047.1 GI:3212944
```

```
KEYWORDS HTG; HTGS_PHASE1.
```

```
SOURCE human.
```

```
ORGANISM Homo sapiens
```

```
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
```

```
REFERENCE 1 (bases 1 to 159103)
```

```
AUTHORS Waterston,R.H.
```

```
TITLE The sequence of Homo sapiens clone
```

```
JOURNAL Unpublished
```

```
REFERENCE 2 (bases 1 to 159103)
```

```
AUTHORS Waterston,R.H.
```

```
TITLE Direct Submission
```

```
JOURNAL Submitted (12-JUN-1998) Genome Sequencing Center, Washington
```

```
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
```

COMMENT

```
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
```

```
* 1 35220: contig of 35220 bp in length
```

```
* 35221 35237: gap of unknown length
```

```
* 35238 159103: contig of 123866 bp in length.
```

```
FEATURES
```

source

```
1..159103
```

```
/organism="Homo sapiens"
```

```
/db_xref="taxon:9606"
```

```
/clone="RG014E15"
```

```
BASE COUNT 49337 a 31915 c 31970 g 45864 t 17 others
```

```
ORIGIN
```

alignment_scores: Quality: 41.00 Length: 11
 Ratio: 4.100 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 72.727

alignment_block:

US-08-653-294-18 x AC005047

Align seg 1/1 to: AC005047 from: 1 to: 159103

1 TyrArgLeuAlaileArgArgileLeuLeuArg 11

62132 TACAGTCTGCAGTAGGAGGAGTCTACTAAAG 62164

seq_name: gb_pr4:AC007687

seq_documentation_block:

LOCUS AC007687 164655 bp DNA PRI 28-OCT-1999
 DEFINITION Homo sapiens 3q26.2-27 BAC RPC111-419H14 (Roswell Park Cancer
 Institute Human BAC Library) complete sequence.

ACCESSION AC007687

VERSION AC007687.16 GI:6137876

KEYWORDS HTG.

SOURCE human.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 164655)
 Muzny,D.M., Adams,C., Bailey,M., Barberia,J., Blankenburg,K.,
 Bodett,B., Bouck,J., Bowie,S., Brooks,A., Buhay,C., Bunac,C.,
 Burkett,C., Burrows,J., Carter,M., Chacko,J., Chen,Z., Cox,C.,
 David,R., Delgado,O., Deshazo,D., Ding,I., Chan,Rashid,N.,
 Durum-Rocha,S., Durbin,K.J., Fernandez,C., Ferraguto,D.,
 Fugum-Tansey,J., Frantz,P., Ganesh,R., Garcia,D.K., Gorrell,J.H.,
 Gorrell,L.L., Guevara,W., Harris,K., He,X., Hernandez,J.,
 Hodgson,A., Hoques,M., Holloway,C., Hosak,H., Jackson,L.E.,
 Jackson,L., Jia,Y., Jones,M., Kelly,S., Kondejewski,N., Kong,Y.,
 Kovar,C., Leal,B., Li,Z., Lichtarge,O., Liu,J., Liu,W., Logan,O.,
 Lu,J., Lucier,R., Martin,R., Martinez,C., McLeod,M.P., Mei,G.,
 Moore,S., Moorish,T., Morgan,M., Morris,S., Nash,S., Nelson,A.,
 Nguyen,R., Nguyen,N., Pu,L.L., Quiles,M., Reiter,D., Rives,M.,
 Payton,B., Perez,L., Pu,L.L., Quiles,M., Reiter,D., Rives,M.,
 Samuel,S., Say,J., Scherer,S., Shah,E., Shen,H., Simon,M.,
 Sparks,A., Stamps,A., Suckang,R., Tabor,P., Taylor,T., Vasquez,L.,
 Vinson,R., Vo,Q., Wahbah,M., Watlington,S., Weinstein,G.,
 Weinstein,I.R., Williamson,A., Worley,K., Wren,J., Wrensford,G.,
 Yu,W., Zhou,X., Naylor,S.L., Nelson,D. and Gibbs,R.

Direct Submission

Unpublished
 2 (bases 1 to 164655)

Worley,K.C.

Direct Submission

Submitted (01-JUN-1999) Molecular and Human Genetics, Baylor
 College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
 3 (bases 1 to 164655)

Worley,K.C.

Direct Submission

Submitted (28-OCT-1999) Human Genome Sequencing Center, Department
 of Molecular and Human Genetics, Baylor College of Medicine, One
 Baylor Plaza, Houston, TX 77030, USA

On Oct 28, 1999 this sequence version replaced gi:6087894.

INFORMATION: http://www.hgsc.bcm.tmc.edu/ or email

gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the
 entire insert of this clone. Overlapping regions of clones are only
 sequenced and submitted once, so the sequence for the remainder of
 the insert may be found in the record for the adjacent clones.
 Overlapping clones are noted at the beginning and end of the
 Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches
 of a local database that includes entries from dbSTS, GDB, and
 local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green,
 unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST
 (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the
 EST and CDNA sequences. Genes demonstrate at least two exons
 flanked by consensus splice sites that maintained sequence
 continuity across the splice junctions. Sequences that are not
 identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum
 standard of double strand coverage with a minimum of 2 clones and 2
 reads with no ambiguities or 2 chemistries with a minimum of 2
 clones and 3 reads with no ambiguities. If the sequence quality for
 a region does not meet this standard, it will be indicated in the
 annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality
 standards - estimated error rate less than 1 per 10,000 bases.
 Reports of lowest quality individual bases and measures of base
 quality are listed below. Description of the metrics can be found
 at URL: <http://gc.bcm.tmc.edu:8088/quality.info/genbank.annotation.html>.

QUALSTAT-REPORT-----

----- Summary Statistics -----
 Contig length: 138071
 Phrap values in estimate: 137060
 Average error rate (BCM-Phrap estimate): 0.000420196
 Fraction of Phrap values less than 40 : 0.036319
 Number of consensus changing edits: 85
 Number of N's in consensus : 0

----- Consensus changing edits -----

Position	Original+Context	Edited+Context
22	gtcggaggac(c)taataacttc	gtcggaggac(n)nnnnnnnnnn
23	tcgaggagac(t)ataaacttcg	tcgaggagac(n)nnnnnnnnnn
24	cgaggagac(t)ataaacttcg	cgaggagac(n)nnnnnnnnnn
25	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
26	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
27	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
28	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
29	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
30	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
31	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
32	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
33	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
34	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
35	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
36	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
37	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
38	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
39	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
40	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
41	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
42	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
43	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
44	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
45	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
46	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
47	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
48	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
49	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
50	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
51	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
52	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn
53	gaggagac(t)ataaacttcg	gaggagac(n)nnnnnnnnnn

1132492 CACAGGCTTGCTGAAGGATTATTTAGTGAGGTAT 132527

CC 1045, 1261, 1483 and 1705).
SQ Sequence 3279 BP; 1505 A; 481 C; 625 G; 668 T;

alignment_scores:
Quality: 37.00 Length: 12
Ratio: 3.083 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 50.000

alignment_block:

US-08-653-294-18 x Q50946/rev ..

Align seg 1/1 to reverse of: Q50946 from: 1 to: 3279

1 TvrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12

2746 TTTCGCTGGTTTTCAGCGTGTGCTTTTAAAGATT 2711

seq_name: N_Geneseq_36:Q51556

seq_documentation_block:

ID Q51556 standard; CDNA; 3279 BP.

AC Q51556;

DE 18-MAY-1994 (first entry)

KW protein; immunoglobulin; binding; immobilisation; light chains;

OS Peptococcus magnus.

FT Key Location/Qualifiers

FT cds 103..3185

FT /tag= a

FT /product= Protein L.

FT repeat_unit 490..573

FT /tag= b

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide positions 673 and 856"

FT repeat_unit 574..672

FT /tag= c

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide position 757"

FT repeat_unit 949..1044

FT /tag= d

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide positions 1162, 1375

FT repeat_unit 1045..1158

FT /tag= e

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide positions 1261, 1483

FT repeat_unit 1822..1938

FT /tag= f

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide positions 2347 and 2545"

FT repeat_unit 1939..2007

FT /tag= g

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide positions 2479, 2665

FT repeat_unit 2035..2094

FT /tag= h

FT /note= "Repeat units are not adjacent, repetitions

of this sequence are not 100% homologous and

begin at nucleotide position 2209"

FT repeat_unit 2095..2208

FT /tag= i

FT /note= "Repeat units are not adjacent, repetitions

FT of this sequence are not 100% homologous and
FT begin at nucleotide positions 2269"
FT 2914..2934
FT /tag= j
FT /note= "Repeat units are adjacent, repetitions
FT of this sequence are not 100% homologous and
FT begin at nucleotide positions 2935, 2953,
FT 2968, 2986, 3001, 3019 and 3034"

PN W09322438-A.

PD 11-NOV-1993.

PF 07-MAY-1993; G00949.

PR 07-MAY-1992; GB-009804.

PA (PUBL-) PUBLIC HEALTH LAB SERVICE BOARD.

PI Atkinson A, Duggleby CJ, Murphy JP, Trowern AR;

DR WPI: 93-368797/46.

DR P-PSDB; R43699.

PT Immunoglobulin binding polypeptide, protein L - used for prodn.

PT of pharmaceuticals and for immobilising antibodies e.g. on

PT columns, in diagnostic tests and in assays

PS Disclosure; Figure 1; 29pp; English.

CC Protein L forms a complex with immunoglobulin kappa light chain.

CC Purified protein can be used as a reagent for immobilising

CC antibodies e.g. on columns, in diagnostic tests and in assays. It

CC may also be used in the production of pharmaceuticals.

SQ Sequence 3279 BP; 1505 A; 480 C; 626 G; 668 T;

alignment_scores:

Quality: 37.00 Length: 12

Ratio: 3.083 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 50.000

alignment_block:

US-08-653-294-18 x Q51556/rev ..

Align seg 1/1 to reverse of: Q51556 from: 1 to: 3279

1 TvrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12

2746 TTTCGCTGGTTTTCAGCGTGTGCTTTTAAAGATT 2711

seq_name: N_Geneseq_36:T17455

seq_documentation_block:

ID T17455 standard; CDNA; 24025 BP.

AC T17455;

DT 07-OCT-1996 (first entry)

DE Mutated BRCA1 genomic sequence from sample set MSKCC family 19921.

KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;

KW antibody production; germline alteration; probe; lesion neoplasia; human;

OS gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.

FT Key Location/Qualifiers

FT exon 256..355

FT /tag= a

FT /note= "exon 1"

FT intron 356..1512

FT /tag= b

FT /note= "intron 1"

FT misc_feature 1295

FT /tag= c

FT /note= "known polymorphic site"

FT exon 1513..1611

FT /tag= d

FT /note= "exon 2"

FT intron 1612..2206

FT /tag= e

FT /note= "intron 2"

FT misc_feature 1925..1937

FT /tag= f

FT /note= "Indefinite interval within intron 2"

FT misc_feature 2141

FT /tag= g

```
FT FT exon /note= "known polymorphic site"
FT FT 2207. .2260
FT FT /tag= h
FT FT /note= "exon 3"
FT FT 2261. .2677
FT FT /tag= i
FT FT /note= "intron 3"
FT FT 2569. .2581
FT FT /tag= j
FT FT /note= "indefinite interval within intron 3"
FT FT 2678. .2788
FT FT /tag= k
FT FT /note= "exon 4"
FT FT 2725
FT FT /tag= l
FT FT /note= "known polymorphic site"
FT FT 2789. .3328
FT FT /tag= m
FT FT /note= "intron 4"
FT FT 3063. .3075
FT FT /tag= n
FT FT /note= "indefinite interval within intron 4"
FT FT 3329. .3406
FT FT /tag= o
FT FT /note= "exon 5"
FT FT 3407. .3813
FT FT /tag= p
FT FT /note= "intron 5"
FT FT 3598. .3610
FT FT /tag= q
FT FT /note= "indefinite interval within intron 5"
FT FT 3653
FT FT /tag= r
FT FT /note= "known polymorphic site"
FT FT 3814. .3902
FT FT /tag= s
FT FT /note= "exon 6"
FT FT 3903. .4224
FT FT /tag= t
FT FT /note= "intron 6"
FT FT 4223
FT FT /tag= u
FT FT /note= "site of 1 nucleotide deletion"
FT FT 4076. .4088
FT FT /tag= v
FT FT /note= "indefinite interval within intron 6"
FT FT 4225. .4364
FT FT /tag= w
FT FT /note= "exon 7"
FT FT 4365. .6571
FT FT /tag= x
FT FT /note= "intron 7"
FT FT 4391. .4392
FT FT /tag= y
FT FT /note= "known polymorphic site"
FT FT 4602. .4614
FT FT /tag= z
FT FT /note= "indefinite interval within intron 7"
FT FT 6538
FT FT /tag= aa
FT FT /note= "known polymorphic site"
FT FT 6572. .6677
FT FT /tag= ab
FT FT /note= "exon 8"
FT FT 6678. .9163
FT FT /tag= ac
FT FT /note= "intron 8"
FT FT 6823
FT FT /tag= ad
FT FT /note= "known polymorphic site"
FT FT 9106
FT FT /tag= ae
FT FT /note= "known polymorphic site"
FT FT 9164. .9209
FT FT /tag= af
FT FT /note= "exon 9"
FT FT 9207
FT FT /tag= ag
FT FT /note= "known polymorphic site"
FT FT 9210. .10530
FT FT /tag= ah
FT FT /note= "intron 9"
FT FT 9376
FT FT /tag= ai
FT FT /note= "known polymorphic site"
FT FT 10531. .10607
FT FT /tag= aj
FT FT /note= "exon 10"
FT FT 10808. .11597
FT FT /tag= ak
FT FT /note= "intron 10"
FT FT 11384. .11396
FT FT /tag= al
FT FT /note= "indefinite interval within intron 10"
FT FT 11598. .15023
FT FT /tag= am
FT FT /note= "exon 11"
FT FT 11908
FT FT /tag= an
FT FT /note= "known polymorphic site"
FT FT 11994
FT FT /tag= ao
FT FT /note= "known polymorphic site"
FT FT 12952
FT FT /tag= ap
FT FT /note= "known polymorphic site"
FT FT 13004
FT FT /tag= aq
FT FT /note= "known polymorphic site"
FT FT 13009
FT FT /tag= ar
FT FT /note= "known polymorphic site"
FT FT 13048
FT FT /tag= as
FT FT /note= "known polymorphic site"
FT FT 13238
FT FT /tag= at
FT FT /note= "known polymorphic site"
FT FT 13448
FT FT /tag= au
FT FT /note= "known polymorphic site"
FT FT 13539
FT FT /tag= av
FT FT /note= "known polymorphic site"
FT FT 13951
FT FT /tag= aw
FT FT /note= "known polymorphic site"
FT FT 14041
FT FT /tag= ax
FT FT /note= "known polymorphic site"
FT FT 14046
FT FT /tag= ay
FT FT /note= "known polymorphic site"
FT FT 14475
FT FT /tag= az
FT FT /note= "known polymorphic site"
FT FT 14874
FT FT /tag= ba
FT FT /note= "known polymorphic site"
FT FT 14891
FT FT /tag= bb
FT FT /note= "known polymorphic site"
FT FT 14966
FT FT /tag= bc
FT FT /note= "known polymorphic site"
FT FT 15024. .15424
FT FT intron
```

```

FT      /tag= bd
FT      /note= "intron 11"
FT      15284
FT      /tag= be
FT      /note= "known polymorphic site"
FT      15425
FT      /tag= bf
FT      /note= "exon 12"
FT      15512
FT      /tag= bg
FT      /note= "intron 12"
FT      15647
FT      /tag= bh
FT      /note= "indefinite interval within intron 12"
FT      15953
FT      /tag= bi
FT      /note= "exon 13"
FT      16077
FT      /tag= bj
FT      /note= "known polymorphic site"
FT      16127
FT      /tag= bk
FT      /note= "intron 13"
FT      16243
FT      /tag= bl

```

```

alignment_scores:
  Quality: 37.00      Length: 11
  Ratio: 3.700      Gaps: 0
  Percent Similarity: 90.909      Percent Identity: 63.636

```

alignment_block:

US-08-653-294-18 x TI7455/rev ..

Align seg 1/1 to reverse of: TI7455 from: 1 to: 24025

```

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||
11313 AGATTGCCATCAGAAACGTGTTCTGATGTAC 11281

```

seq_name: N_Geneseq_36:TI7515

seq_documentation_block:

```

ID   TI7515 standard; cDNA: 24025 BP.
AC   TI7515;
DT   04-OCT-1996 (first entry)
DE   Mutated BRCA1 genomic sequence from PM15.
KW   Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
KW   antibody production; germline alteration; probe; lesion neoplasia; human;
KW   gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
OS   Homo sapiens.
FH   Key
FT      Location/Qualifiers
FT      exon
FT      256..355
FT      /tag= a
FT      /note= "exon 1"
FT      356..1512
FT      /tag= b
FT      /note= "intron 1"
FT      1295
FT      /tag= c
FT      /note= "known polymorphic site"
FT      1513..1611
FT      /tag= d
FT      /note= "exon 2"
FT      1612..2206
FT      /tag= e
FT      /note= "intron 2"
FT      1925..1937
FT      /tag= f
FT      /note= "indefinite interval within intron 2"
FT      2141
FT      /tag= g
FT      /note= "known polymorphic site"

```

```

FT      exon
FT      2207..2260
FT      /tag= h
FT      /note= "exon 3"
FT      2261..2677
FT      /tag= i
FT      /note= "intron 3"
FT      2569..2581
FT      /tag= j
FT      /note= "indefinite interval within intron 3"
FT      2678..2788
FT      /tag= k
FT      /note= "exon 4"
FT      2725
FT      /tag= l
FT      /note= "known polymorphic site"
FT      2789..3328
FT      /tag= m
FT      /note= "intron 4"
FT      3063..3075
FT      /tag= n
FT      /note= "indefinite interval within intron 4"
FT      3329..3406
FT      /tag= o
FT      /note= "exon 5"
FT      3407..3813
FT      /tag= p
FT      /note= "intron 5"
FT      3598..3610
FT      /tag= q
FT      /note= "indefinite interval within intron 5"
FT      3653
FT      /tag= r
FT      /note= "known polymorphic site"
FT      3814..3902
FT      /tag= s
FT      /note= "exon 6"
FT      3903..4224
FT      /tag= t
FT      /note= "intron 6"
FT      4076..4088
FT      /tag= u
FT      /note= "indefinite interval within intron 6"
FT      4225..4364
FT      /tag= v
FT      /note= "exon 7"
FT      4365..6571
FT      /tag= w
FT      /note= "intron 7"
FT      4391..4392
FT      /tag= x
FT      /note= "known polymorphic site"
FT      4602..4614
FT      /tag= y
FT      /note= "indefinite interval within intron 7"
FT      6538
FT      /tag= z
FT      /note= "known polymorphic site"
FT      6572..6677
FT      /tag= aa
FT      /note= "exon 8"
FT      6678..9163
FT      /tag= ab
FT      /note= "intron 8"
FT      6823
FT      /tag= ac
FT      /note= "known polymorphic site"
FT      9106
FT      /tag= ad
FT      /note= "site of 1 nucleotide deletion at known polymorphic site"
FT      9163..9208
FT      /tag= ae
FT      /note= "exon 9"

```

```
FT misc_feature 9206 /*tag= af
FT /note= "known polymorphic site"
FT 9209. .10529
FT /*tag= ag
FT /note= "intron 9"
FT 9375
FT /*tag= ah
FT /note= "known polymorphic site"
FT 10530. .10606
FT /*tag= ai
FT /note= "exon 10"
FT 10607. .11596
FT /*tag= aj
FT /note= "intron 10"
FT 11383. .11395
FT /*tag= ak
FT /note= "indefinite interval within intron 10"
FT 11597. .15022
FT /*tag= al
FT /note= "exon 11"
FT 11907
FT /*tag= am
FT /note= "known polymorphic site"
FT 11993
FT /*tag= an
FT /note= "known polymorphic site"
FT 12951
FT /*tag= ao
FT /note= "known polymorphic site"
FT 13003
FT /*tag= ap
FT /note= "known polymorphic site"
FT 13008
FT /*tag= aq
FT /note= "known polymorphic site"
FT 13047
FT /*tag= ar
FT /note= "known polymorphic site"
FT 13237
FT /*tag= as
FT /note= "known polymorphic site"
FT 13447
FT /*tag= at
FT /note= "known polymorphic site"
FT 13538
FT /*tag= au
FT /note= "known polymorphic site"
FT 13950
FT /*tag= av
FT /note= "known polymorphic site"
FT 14040
FT /*tag= aw
FT /note= "known polymorphic site"
FT 14045
FT /*tag= ax
FT /note= "known polymorphic site"
FT 14474
FT /*tag= ay
FT /note= "known polymorphic site"
FT 14873
FT /*tag= az
FT /note= "known polymorphic site"
FT 14890
FT /*tag= ba
FT /note= "known polymorphic site"
FT 14965
FT /*tag= bb
FT /note= "known polymorphic site"
FT 15023. .15423
FT /*tag= bc
FT /note= "intron 11"
FT 15283
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```
FT exon /*tag= bd
FT /note= "known polymorphic site"
FT 15424. .15510
FT /*tag= be
FT /note= "exon 12"
FT 15511. .15951
FT /*tag= bf
FT /note= "intron 12"
FT 15646. .15658
FT /*tag= bg
FT /note= "indefinite interval within intron 12"
FT 15952. .16125
FT /*tag= bh
FT /note= "exon 13"
FT 16076
FT /*tag= bi
FT /note= "known polymorphic site"
FT 16126. .16564
FT /*tag= bj
FT /note= "intron 13"
FT 16242
FT /*tag= bk
FT /note= "known polymorphic site"
FT 16369. .16381

alignment_scores:
Quality: 37.00 Length: 11
Ratio: 3.700 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
US-08-653-294-18 x TI17515/rev ..
Align seg 1/1 to reverse of: TI17515 from: 1 to: 24025

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||
11313 AGATTGGCCATCAGAAACCTGGTCTGTGATGTAC 11281

seq_name: N_Geneseq_36:TI18325

seq_documentation_block:
ID TI18325 standard; DNA; 24026 BP.
AC TI18325;
DT 05-JUN-1996 (first entry)
DE BRCA1, human breast and ovarian cancer predisposing gene.
KW BRCA1; breast cancer; ovary cancer; predisposing gene;
KW susceptibility gene; diagnosis; prognosis; gene therapy; ds.
OS Homo sapiens.
PH Key Location/Qualifiers
FT intron 1. .55
FT /*tag= a
FT exon 56. .155
FT /*tag= b
FT intron 156. .1512
FT /*tag= c
FT exon 1513. .1611
FT /*tag= d
FT intron 1612. .2206
FT /*tag= e
FT /note= "n at 1925-1937 represent an indefinite
FT interval within the intron"
FT 2207. .2260
FT /*tag= f
FT exon 2261. .2677
FT /*tag= g
FT /note= "n at 2569-2581 represent an indefinite
FT interval within the intron"
FT 2678. .2788
FT /*tag= h
FT intron 2789. .3328
FT /*tag= i
FT /note= "n at 3063-3075 represent an indefinite
```

```
FT FT exon interval within the intron"
FT 3329. .3406
FT /*tag= j
FT intron 3407. .3813
FT /*tag= k
FT /note= "n at 3598-3610 represent an indefinite
FT interval within the intron"
FT 3814. .3902
FT /*tag= l
FT intron 3903. .4224
FT /*tag= m
FT /note= "n at 4076-4088 represent an indefinite
FT interval within the intron"
FT 4225. .4364
FT /*tag= n
FT intron 4365. .6571
FT /*tag= o
FT /note= "n at 4602-4614 represent an indefinite
FT interval within the intron"
FT 6572. .6677
FT /*tag= p
FT intron 6678. .9163
FT /*tag= q
FT exon 9164. .9207
FT /*tag= r
FT intron 9208. .10530
FT /*tag= s
FT exon 10531. .10607
FT /*tag= t
FT intron 10608. .11597
FT /*tag= u
FT /note= "n at 11383-11396 represent an indefinite
FT interval within the intron"
FT 11598. .15023
FT /*tag= v
FT intron 15024. .15424
FT /*tag= w
FT exon 15425. .15511
FT /*tag= x
FT intron 15512. .15952
FT /*tag= y
FT /note= "n at 15647-15659 represent an indefinite
FT interval within the intron"
FT 15953. .16126
FT /*tag= z
FT intron 16127. .16565
FT /*tag= aa
FT /note= "n at 16370-16382 represent an indefinite
FT interval within the intron"
FT 16566. .16692
FT /*tag= ab
FT intron 16693. .17535
FT /*tag= ac
FT /note= "n at 17390-17302 represent an indefinite
FT interval within the intron"
FT 17536. .17726
FT /*tag= ad
FT intron 17727. .18416
FT /*tag= ae
FT /note= "n at 18399-18312 represent an indefinite
FT interval within the intron"
FT 18417. .18787
FT /*tag= af
FT intron 18788. .19298
FT /*tag= ag
FT /note= "n at 18952-18964 represent an indefinite
FT interval within the intron"
FT 19299. .19386
FT /*tag= ah
FT exon 19387. .20190
FT /*tag= ai
FT /note= "n at 19887-19899 represent an indefinite
FT interval within the intron"
FT 20191. .20267
FT /*tag= aj
FT intron 20268. .21094
FT /*tag= ak
FT /note= "n at 20767-20779 represent an indefinite
FT interval within the intron"
FT 21095. .21135
FT /*tag= al
FT intron 21136. .21583
FT /*tag= am
FT /note= "n at 21341-21353 represent an indefinite
FT interval within the intron"
FT 21584. .21667
FT /*tag= an
FT intron 21668. .22233
FT /*tag= ao
FT /note= "n at 21921-21933 represent an indefinite
FT interval within the intron"
FT 22234. .22288
FT /*tag= ap
FT intron 22289. .22832
FT /*tag= aq
FT /note= "n at 22567-22579 represent an indefinite
FT interval within the intron"
FT 22833. .22906
FT /*tag= ar
FT exon 22907. .23287
FT /*tag= as
FT intron 23288. .23348
FT /*tag= at
FT exon 23349. .23698
FT /*tag= au
FT /note= "n at 23580-23592 represent an indefinite
FT interval within the intron"
FT 23699. .24026
FT /*tag= av
FT exon 2725
FT misc_feature /*tag= aw
FT /note= "polymorphic site"
FT 3653
FT misc_feature /*tag= ax
FT /note= "polymorphic site"
FT 4391
FT misc_feature /*tag= ay
FT /note= "polymorphic site"
FT 4392
FT misc_feature /*tag= az
FT /note= "polymorphic site"
FT 6823
FT misc_feature /*tag= ba
FT /note= "polymorphic site"
FT 9106
FT misc_feature /*tag= bb
FT /note= "polymorphic site"
FT 9207
FT misc_feature /*tag= bc
FT /note= "polymorphic site"
FT 9376
FT misc_feature /*tag= bd
FT /note= "polymorphic site"
FT 11908
FT misc_feature /*tag= be
FT /note= "polymorphic site"
FT 11994
FT misc_feature /*tag= bf
FT /note= "polymorphic site"
FT 12952
FT misc_feature /*tag= bg
FT /note= "polymorphic site"
FT 13004
FT misc_feature /*tag= bh
```



```

FT      /note= "polymorphic site"
FT      13009
FT      /tag= bi
FT      /note= "polymorphic site"
FT      13048
FT      /tag= bj
FT      /note= "polymorphic site"
FT      13238
FT      /tag= bk
FT      /note= "polymorphic site"
FT      13448
FT      /tag= bl
FT      /note= "polymorphic site"
FT      13539
FT      /tag= bm
FT      /note= "polymorphic site"
FT      13951
FT      /tag= bn
FT      /note= "polymorphic site"
FT      14041
FT      /tag= bo
FT      /note= "polymorphic site"
FT      14046

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alignment_scores: Quality: 37.00 Length: 11
 Ratio: 3.700 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

us-08-653-294-18 x T18325/rev ..
 Align seg 1/1 to reverse of: T18325 from: 1 to: 24026

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12

|||||
 11314 AGATTGCCATCAGAAACGTGCTGTGATAC 11282

seq_name: N_Geneseq_36:T17512

seq_documentation_block:

ID T17512 standard; cDNA: 24026 BP.
 AC T17512;
 DT 04-OCT-1996 (first entry)
 DE Mutated BRCA1 genomic sequence from PM04.
 KW Cancer therapy; breast and ovarian cancer
 KW antibody production; germline alteration; probe; lesion neoplasia; human;
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT 256..355
 FT /tag= a
 FT /note= "exon 1"
 FT 356..1512
 FT /tag= b
 FT /note= "intron 1"
 FT 1295
 FT /tag= c
 FT /note= "known polymorphic site"
 FT 1513..1611
 FT /tag= d
 FT /note= "exon 2"
 FT 1612..2206
 FT /tag= e
 FT /note= "intron 2"
 FT 1925..1937
 FT /tag= f
 FT /note= "indefinite interval within intron 2"
 FT 2141
 FT /tag= g
 FT /note= "known polymorphic site"
 FT 2207..2260
 FT /tag= h

```

FT      intron
FT      2261..2677
FT      /tag= i
FT      /note= "intron 3"
FT      2569..2581
FT      /tag= j
FT      /note= "indefinite interval within intron 3"
FT      2678..2788
FT      /tag= k
FT      /note= "exon 4"
FT      2725
FT      /tag= l
FT      /note= "known polymorphic site"
FT      2789..3328
FT      /tag= m
FT      /note= "intron 4"
FT      3063..3075
FT      /tag= n
FT      /note= "indefinite interval within intron 4"
FT      3329..3406
FT      /tag= o
FT      /note= "exon 5"
FT      3407..3813
FT      /tag= p
FT      /note= "intron 5"
FT      3598..3610
FT      /tag= q
FT      /note= "indefinite interval within intron 5"
FT      3653
FT      /tag= r
FT      /note= "known polymorphic site"
FT      3814..3902
FT      /tag= s
FT      /note= "exon 6"
FT      3903..4224
FT      /tag= t
FT      /note= "intron 6"
FT      4076..4088
FT      /tag= u
FT      /note= "indefinite interval within intron 6"
FT      4225..4364
FT      /tag= v
FT      /note= "exon 7"
FT      4365..6571
FT      /tag= w
FT      /note= "intron 7"
FT      4391..4392
FT      /tag= x
FT      /note= "known polymorphic site"
FT      4602..4614
FT      /tag= y
FT      /note= "indefinite interval within intron 7"
FT      6538
FT      /tag= z
FT      /note= "known polymorphic site"
FT      6572..6677
FT      /tag= aa
FT      /note= "exon 8"
FT      6678..9163
FT      /tag= ab
FT      /note= "intron 8"
FT      6823
FT      /tag= ac
FT      /note= "known polymorphic site"
FT      9106
FT      /tag= ad
FT      /note= "known polymorphic site"
FT      9164..9209
FT      /tag= ae
FT      /note= "exon 9"
FT      9207
FT      /tag= af
FT      /note= "known polymorphic site"

```

```
FT intron 9210..10530
FT /*tag= ag
FT /note= "intron 9"
FT 9376
FT /*tag= ah
FT /note= "known polymorphic site"
FT 10531..10607
FT /*tag= ai
FT /note= "exon 10"
FT 10808..11597
FT /*tag= aj
FT /note= "intron 10"
FT 11384..11396
FT /*tag= ak
FT /note= "indefinite interval within intron 10"
FT 11598..15023
FT /*tag= al
FT /note= "exon 11"
FT 11908
FT /*tag= am
FT /note= "known polymorphic site"
FT 11994
FT /*tag= an
FT /note= "known polymorphic site"
FT 12952
FT /*tag= ao
FT /note= "known polymorphic site"
FT 13004
FT /*tag= ap
FT /note= "known polymorphic site"
FT 13009
FT /*tag= ag
FT /note= "known polymorphic site"
FT 13048
FT /*tag= ar
FT /note= "known polymorphic site"
FT 13238
FT /*tag= as
FT /note= "known polymorphic site"
FT 13448
FT /*tag= at
FT /note= "known polymorphic site"
FT 13539
FT /*tag= au
FT /note= "known polymorphic site"
FT 13951
FT /*tag= av
FT /note= "known polymorphic site"
FT 14041
FT /*tag= aw
FT /note= "known polymorphic site"
FT 14046
FT /*tag= ax
FT /note= "known polymorphic site"
FT 14475
FT /*tag= ay
FT /note= "known polymorphic site"
FT 14874
FT /*tag= az
FT /note= "known polymorphic site"
FT 14891
FT /*tag= ba
FT /note= "known polymorphic site"
FT 14966
FT /*tag= bb
FT /note= "known polymorphic site"
FT 15024..15424
FT /*tag= bc
FT /note= "intron 11"
FT 15284
FT /*tag= bd
FT /note= "C to A mutation at known polymorphic site"
FT 15425..15511
```

```
FT /*tag= be
FT /note= "exon 12"
FT 15512..15952
FT /*tag= bf
FT /note= "intron 12"
FT 15647..15659
FT /*tag= bg
FT /note= "indefinite interval within intron 12"
FT 15953..16126
FT /*tag= bh
FT /note= "exon 13"
FT 16077
FT /*tag= bi
FT /note= "known polymorphic site"
FT 16127..16565
FT /*tag= bj
FT /note= "intron 13"
FT 16243
FT /*tag= bk
FT /note= "known polymorphic site"
FT 16370..16382
FT /*tag= bl

alignment_scores:
  Quality: 37.00 Length: 11
  Ratio: 3.700 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
US-08-653-294-18 x T17512/rev ..
Align seg 1/1 to reverse of: T17512 from: 1 to: 24026

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||
11314 AGATTGCCATCAGAAACTGGTTCTGATGTAC 11282

seq_name: N_Geneseq_36:T17513

seq_documentation_block:
ID T17513 standard; CDNA; 24026 BP.
AC T17513;
DT 04-OCT-1996 (first entry)
DE Mutated BRCA1 genomic sequence from PM05.
KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
KW antibody production; germline alteration; probe; lesion neoplasia; human;
KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
OS Homo sapiens.
FH Key Location/Qualifiers
FT exon 256..355
FT /*tag= a
FT /note= "exon 1"
FT 356..1512
FT /*tag= b
FT /note= "intron 1"
FT 1295
FT misc_feature
FT /*tag= c
FT /note= "known polymorphic site"
FT 1513..1611
FT /*tag= d
FT /note= "exon 2"
FT 1612..2206
FT /*tag= e
FT /note= "intron 2"
FT 1925..1937
FT misc_feature
FT /*tag= f
FT /note= "indefinite interval within intron 2"
FT 2141
FT misc_feature
FT /*tag= g
FT /note= "known polymorphic site"
FT 2207..2260
FT /*tag= h
FT /note= "exon 3"
```

```
FT intron 2261..2677 /*tag= i /*tag= ag
FT /note= "intron 3" /note= "intron 9"
FT misc_feature 9376
FT exon /*tag= ah /*tag= ah
FT /*note= "known polymorphic site" /*note= "known polymorphic site"
FT 10531..10607 /*tag= ai
FT /*note= "exon 10" /*note= "exon 10"
FT 10608..11597 /*tag= aj
FT /*note= "intron 10" /*note= "intron 10"
FT 11384..11396 /*tag= ak
FT /*note= "indefinite interval within intron 10"
FT exon 11598..15023 /*tag= al
FT /*note= "exon 11"
FT misc_feature 11908
FT /*tag= am
FT /*note= "known polymorphic site"
FT misc_feature 11994
FT /*tag= an
FT /*note= "known polymorphic site"
FT misc_feature 12952
FT /*tag= ao
FT /*note= "known polymorphic site"
FT misc_feature 13004
FT /*tag= ap
FT /*note= "known polymorphic site"
FT misc_feature 13009
FT /*tag= aq
FT /*note= "known polymorphic site"
FT misc_feature 13048
FT /*tag= ar
FT /*note= "known polymorphic site"
FT misc_feature 13238
FT /*tag= as
FT /*note= "known polymorphic site"
FT misc_feature 13448
FT /*tag= at
FT /*note= "known polymorphic site"
FT misc_feature 13539
FT /*tag= au
FT /*note= "known polymorphic site"
FT misc_feature 13951
FT /*tag= av
FT /*note= "known polymorphic site"
FT misc_feature 14041
FT /*tag= aw
FT /*note= "known polymorphic site"
FT misc_feature 14046
FT /*tag= ax
FT /*note= "known polymorphic site"
FT misc_feature 14475
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FT misc_feature 14891
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FT /*note= "known polymorphic site"
FT intron 15024..15424 /*tag= bc
FT /*note= "intron 11"
FT misc_feature 15284
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FT exon 15425..15511 /*tag= be
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FT misc_feature 9376
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FT /*note= "known polymorphic site" /*note= "known polymorphic site"
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FT /*note= "exon 10" /*note= "exon 10"
FT 10608..11597 /*tag= aj
FT /*note= "intron 10" /*note= "intron 10"
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FT /*note= "indefinite interval within intron 10"
FT exon 11598..15023 /*tag= al
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FT misc_feature 11908
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FT /*note= "known polymorphic site"
FT misc_feature 11994
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FT /*note= "intron 11"
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FT exon 15425..15511 /*tag= be
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FT		/note= "intron 4"	
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FT		/*tag= q	
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FT	intron	3903. .4224	
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FT		/note= "intron 6"	
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FT	exon	4225. .4364	
FT		/*tag= v	
FT		/note= "exon 7"	
FT	intron	4365. .6571	
FT		/*tag= w	
FT		/note= "intron 7"	
FT	misc_feature	4391. .4392	
FT		/*tag= x	
FT		/note= "known polymorphic site"	
FT	misc_feature	4602. .4614	
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FT		/note= "indefinite interval within intron 7"	
FT	misc_feature	6338	
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FT	intron	6678. .9163	
FT		/*tag= ab	
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FT		/*tag= ac	
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FT	misc_feature	9106	
FT		/*tag= ad	
FT		/note= "known polymorphic site"	
FT	exon	9164. .9209	
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FT		/note= "exon 9"	
FT	misc_feature	9207	
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FT		/note= "known polymorphic site"	
FT	intron	9210. .10530	
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FT FT misc_feature /note= "intron 9"
FT FT 9376 /tag= ah
FT FT /note= "known polymorphic site"
FT FT 10531.10607 /tag= ai
FT FT /note= "exon 10"
FT FT 10608.11597 /tag= aj
FT FT /note= "intron 10"
FT FT 11384.11396 /tag= ak
FT FT /note= "indefinite interval within intron 10"
FT FT 11598.15023 /tag= al
FT FT /note= "exon 11"
FT FT 11908 /tag= am
FT FT /note= "known polymorphic site"
FT FT 11994 /tag= an
FT FT /note= "known polymorphic site"
FT FT 12952 /tag= ao
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FT FT /note= "known polymorphic site"
FT FT 13539 /tag= au
FT FT /note= "known polymorphic site"
FT FT 13951 /tag= av
FT FT /note= "known polymorphic site"
FT FT 14041 /tag= aw
FT FT /note= "known polymorphic site"
FT FT 14046 /tag= ax
FT FT /note= "known polymorphic site"
FT FT 14475 /tag= ay
FT FT /note= "known polymorphic site"
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FT FT /note= "known polymorphic site"
FT FT 14891 /tag= ba
FT FT /note= "known polymorphic site"
FT FT 14966 /tag= bb
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FT FT 15024.15424 /tag= bc
FT FT /note= "intron 11"
FT FT 15284 /tag= bd
FT FT /note= "known polymorphic site"
FT FT 15425.15511 /tag= be
FT FT /note= "exon 12"
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FT FT intron 15512.15952 /tag= bf
FT FT /note= "intron 12"
FT FT 15847.15659 /tag= bg
FT FT /note= "indefinite interval within intron 12"
FT FT 15953.16126 /tag= bh
FT FT /note= "exon 13"
FT FT 16077 /tag= bi
FT FT /note= "known polymorphic site"
FT FT 16127.16565 /tag= bj
FT FT /note= "intron 13"
FT FT 16243 /tag= bk
FT FT /note= "known polymorphic site"
FT FT 16370.16382 /tag= bl
FT FT /note= "intron 13"

alignment_scores:
  Quality: 37.00 Length: 11
  Ratio: 3.700 Gaps: 0
  Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
  US-08-653-294-18 x T17514/rev ..
  Align seg 1/1 to reverse of: T17514 from: 1 to: 24026
  2 ArgLeuAlaIleArgIleLeuLeuArgTyr 12
  |||||
  11314 AGATTGGCCATCAGAAACTGGTCTGATGTAC 11282

seq_name: N_Geneseq_36:T17516

seq_documentation_block:
  ID T17516 standard; cDNA; 24026 BP.
  AC T17516;
  DT 04-OCT-1996 (first entry)
  DE Mutated BRCA1 genomic sequence from PM16.
  KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
  KW antibody production; germline alteration; probe; lesion neoplasia; human;
  KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
  OS Homo sapiens.
  FH key Location/Qualifiers
  FT FT exon 256..355
  FT FT /tag= a
  FT FT /note= "exon 1"
  FT FT 356..1512 /tag= b
  FT FT /note= "intron 1"
  FT FT 1295 /tag= c
  FT FT /note= "known polymorphic site"
  FT FT 1513..1611 /tag= d
  FT FT /note= "exon 2"
  FT FT 1612..2206 /tag= e
  FT FT /note= "intron 2"
  FT FT 1925..1937 /tag= f
  FT FT /note= "indefinite interval within intron 2"
  FT FT 2141 /tag= g
  FT FT /note= "known polymorphic site"
  FT FT 2207..2260 /tag= h
  FT FT /note= "exon 3"
  FT FT 2261..2677 /tag= i
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FT 2725
FT /tag= l
FT /note= "known polymorphic site"
FT 2789. .3328
FT /tag= m
FT /note= "intron 4"
FT 3063. .3075
FT /tag= n
FT /note= "indefinite interval within intron 4"
FT 3329. .3406
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FT /note= "exon 5"
FT 3407. .3813
FT /tag= p
FT /note= "intron 5"
FT 3598. .3610
FT /tag= q
FT /note= "indefinite interval within intron 5"
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FT /note= "known polymorphic site"
FT 3814. .3902
FT /tag= s
FT /note= "exon 6"
FT 3903. .4224
FT /tag= t
FT /note= "intron 6"
FT 4076. .4088
FT /tag= u
FT /note= "indefinite interval within intron 6"
FT 4225. .4364
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FT /note= "exon 7"
FT 4365. .6571
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FT /note= "known polymorphic site"
FT 4602. .4614
FT /tag= y
FT /note= "indefinite interval within intron 7"
FT 6538
FT /tag= z
FT /note= "known polymorphic site"
FT 6572. .6677
FT /tag= aa
FT /note= "exon 8"
FT 6678. .9163
FT /tag= ab
FT /note= "intron 8"
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FT 9164. .9209
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FT /note= "exon 9"
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FT 9210. .10530
FT /tag= ag
FT /note= "intron 9"
FT 9376
FT /tag= ah
FT /note= "known polymorphic site"
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FT /tag= ai
FT /note= "exon 10"
FT 10608. .11597
FT /tag= aj
FT /note= "intron 10"
FT 11384. .11396
FT /tag= ak
FT /note= "indefinite interval within intron 10"
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FT /tag= al
FT /note= "exon 11"
FT 11908
FT /tag= am
FT /note= "known polymorphic site"
FT 11994
FT /tag= an
FT /note= "known polymorphic site"
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FT 13009
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FT 13048
FT /tag= ar
FT /note= "known polymorphic site"
FT 13238
FT /tag= as
FT /note= "known polymorphic site"
FT 13448
FT /tag= at
FT /note= "known polymorphic site"
FT 13539
FT /tag= au
FT /note= "known polymorphic site"
FT 13951
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FT /note= "known polymorphic site"
FT 14041
FT /tag= aw
FT /note= "known polymorphic site"
FT 14046
FT /tag= ax
FT /note= "known polymorphic site"
FT 14475
FT /tag= ay
FT /note= "known polymorphic site"
FT 14874
FT /tag= az
FT /note= "known polymorphic site"
FT 14891
FT /tag= ba
FT /note= "known polymorphic site"
FT 14966
FT /tag= bb
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FT 15024. .15424
FT /tag= bc
FT /note= "intron 11"
FT 15284
FT /tag= bd
FT /note= "known polymorphic site"
FT 15425. .15511
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FT      /tag= bi
FT      /note= "known polymorphic site"
FT      /tag= bj
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alignment_scores:
 Quality: 37.00 Length: 11
 Ratio: 3.700 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-08-653-294-18 x TL7516/rev ..
 Align seg 1/1 to reverse of: TL7516 from: 1 to: 24026

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
 11314 AGATTGCCATCAAGAAACGGTTCTGATGAC 11282

seq_name: N_Geneseq_36:TL7517

seq_documentation_block:

ID TL7517 standard; CDNA: 24026 BP.
 AC TL7517;
 DT 04-OCT-1996 (first entry)
 DE Mutated BRCA1 genomic sequence from PMA02.1.
 KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
 KW antibody production; germline alteration; probe; lesion neoplasia; human;
 KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
 OS Homo sapiens.
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 FT 256..355
 FT /tag= a
 FT /note= "exon 1"
 FT 356..1512
 FT /tag= b
 FT /note= "intron 1"
 FT 1295
 FT /tag= c
 FT /note= "G to A mutation at known polymorphic site"
 FT 1513..1611
 FT /tag= d
 FT /note= "exon 2"
 FT 1612..2206
 FT /tag= e
 FT /note= "intron 2"
 FT 1925..1937
 FT /tag= f
 FT /note= "indefinite interval within intron 2"
 FT 2141
 FT /tag= g
 FT /note= "known polymorphic site"
 FT 2207..2260
 FT /tag= h
 FT /note= "exon 3"
 FT 2261..2677
 FT /tag= i
 FT /note= "intron 3"

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FT      2678..2788
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FT      /note= "exon 4"
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FT      2789..3328
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FT      /note= "intron 4"
FT      3063..3075
FT      /tag= n
FT      /note= "indefinite interval within intron 4"
FT      3329..3406
FT      /tag= o
FT      /note= "exon 5"
FT      3407..3813
FT      /tag= p
FT      /note= "intron 5"
FT      3598..3610
FT      /tag= q
FT      /note= "indefinite interval within intron 5"
FT      3653
FT      /tag= r
FT      /note= "known polymorphic site"
FT      3814..3902
FT      /tag= s
FT      /note= "exon 6"
FT      3903..4224
FT      /tag= t
FT      /note= "intron 6"
FT      4076..4088
FT      /tag= u
FT      /note= "indefinite interval within intron 6"
FT      4225..4364
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FT      4365..6571
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FT      4391..4392
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FT      /note= "known polymorphic site"
FT      4602..4614
FT      /tag= y
FT      /note= "indefinite interval within intron 7"
FT      6538
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FT      6572..6677
FT      /tag= aa
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FT      6678..9163
FT      /tag= ab
FT      /note= "intron 8"
FT      6823
FT      /tag= ac
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FT      9106
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FT      /note= "known polymorphic site"
FT      9164..9209
FT      /tag= ae
FT      /note= "exon 9"
FT      9207
FT      /tag= af
FT      /note= "known polymorphic site"
FT      9210..10530
FT      /tag= ag
FT      /note= "intron 9"
FT      9376

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FT /*tag= "exon 10"
FT intron 10508. .11597 /*tag= aj
FT /*tag= "intron 10"
FT misc_feature 11384. .11396 /*tag= ak
FT /*tag= "indefinite interval within intron 10"
FT exon 11598. .15023 /*tag= al
FT /*tag= "exon 11"
FT misc_feature 11908 /*tag= am
FT /*tag= "known polymorphic site"
FT misc_feature 11994 /*tag= an
FT /*tag= "known polymorphic site"
FT misc_feature 12952 /*tag= ao
FT /*tag= "known polymorphic site"
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FT misc_feature 13009 /*tag= aq
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FT /*tag= "known polymorphic site"
FT misc_feature 14966 /*tag= bb
FT /*tag= "known polymorphic site"
FT intron 15024. .15424 /*tag= bc
FT /*tag= "intron 11"
FT misc_feature 15284 /*tag= bd
FT /*tag= "known polymorphic site"
FT exon 15425. .15511 /*tag= be
FT /*tag= "exon 12"
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FT FT /*note= "intron 12"
FT misc_feature 15647. .15659 /*tag= bg
FT /*tag= "indefinite interval within intron 12"
FT exon 15953. .16126 /*tag= bh
FT /*tag= "exon 13"
FT misc_feature 16077 /*tag= bi
FT /*tag= "known polymorphic site"
FT intron 16127. .16565 /*tag= bj
FT /*tag= "intron 13"
FT misc_feature 16243 /*tag= bk
FT /*tag= "known polymorphic site"
FT misc_feature 16370. .16382 /*tag= bl

alignment_scores:
  Quality: 37.00 Length: 11
  Ratio: 3.700 Gaps: 0
  Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
  US-08-653-294-18 x T17517/rev ..
  Align seg 1/1 to reverse of: T17517 from: 1 to: 24026

      2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
      |||||
11314 AGATTGCCATCAGAAACTGGTCTGATGTAC 11282

seq_name: N_Geneseq_36:T17518

seq_documentation_block:
ID T17518 standard; cDNA; 24026 BP.
AC T17518;
DT 04-OCT-1996 (first entry)
DE Mutated BRCA1 genomic sequence from PMA03.1.
KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
KW antibody production; germline alteration; probe; lesion neoplasia; human;
KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
OS Homo sapiens.
FH Key
FT exon 256. .355 Location/Qualifiers
FT /*tag= a /note= "exon 1"
FT /*tag= 356. .1512
FT intron
FT /*tag= b /note= "intron 1"
FT misc_feature 1295
FT /*tag= c /note= "known polymorphic site"
FT exon 1513. .1611
FT /*tag= d /note= "exon 2"
FT intron 1612. .2206
FT /*tag= e /note= "intron 2"
FT misc_feature 1925. .1937
FT /*tag= f /note= "indefinite interval within intron 2"
FT mutation 2141
FT /*tag= g /note= "G to C mutation at known polymorphic site"
FT exon 2207. .2260
FT /*tag= h /note= "exon 3"
FT intron 2261. .2677
FT /*tag= i /note= "intron 3"
FT misc_feature 2569. .2581
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FT misc_feature 15647..15659      /tag= bg
FT /note= "indefinite interval within intron 12"
FT exon 15953..16126
FT /tag= bh
FT /note= "exon 13"
FT misc_feature 16077
FT /tag= bi
FT /note= "known polymorphic site"
FT intron 16127..16565
FT /tag= bj
FT /note= "intron 13"
FT misc_feature 16243
FT /tag= bk
FT /note= "known polymorphic site"
FT misc_feature 16370..16382
FT /tag= bl

alignment_scores:
  Quality: 37.00      Length: 11
  Ratio: 3.700      Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
US-08-653-294-18 x TI7518/rev ..
Align seg 1/1 to reverse of: TI7518 from: 1 to: 24026

      2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
      11314 AGATTGCCCATCAGAAACTGTTCTGTGATGAC 11282

seq_name: N_Geneseq_36:TI7519
seq_documentation_block:
ID TI7519 standard; cDNA; 24026 BP.
AC TI7519;
DE 04-OCT-1996 (first entry)
DE Mutated BRCA1 genomic sequence from PMA06.1.
KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
KW antibody production; germline alteration; probe; lesion neoplasia; human;
KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
OS Homo sapiens.
FH Key
FT 256..355      Location/Qualifiers
FT /tag= a
FT /note= "exon 1"
FT intron 356..1512
FT /tag= b
FT /note= "intron 1"
FT misc_feature 1295
FT /tag= c
FT /note= "known polymorphic site"
FT exon 1513..1611
FT /tag= d
FT /note= "exon 2"
FT intron 1612..2206
FT /tag= e
FT /note= "intron 2"
FT misc_feature 1925..1937
FT /tag= f
FT /note= "indefinite interval within intron 2"
FT misc_feature 2141
FT /tag= g
FT /note= "known polymorphic site"
FT exon 2207..2260
FT /tag= h
FT /note= "exon 3"
FT intron 2261..2677
FT /tag= i
FT /note= "intron 3"
FT misc_feature 2569..2581
FT /tag= j
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FT 2678..2788
FT /tag= k
FT /note= "exon 4"
FT misc_feature 2725
FT /tag= l
FT /note= "known polymorphic site"
FT intron 2789..3328
FT /tag= m
FT /note= "intron 4"
FT misc_feature 3063..3075
FT /tag= n
FT /note= "indefinite interval within intron 4"
FT exon 3329..3406
FT /tag= o
FT /note= "exon 5"
FT intron 3407..3813
FT /tag= p
FT /note= "intron 5"
FT misc_feature 3598..3610
FT /tag= q
FT /note= "indefinite interval within intron 5"
FT mutation 3653
FT /tag= r
FT /note= "A to G mutation at known polymorphic site"
FT exon 3814..3902
FT /tag= s
FT /note= "exon 6"
FT intron 3903..4224
FT /tag= t
FT /note= "intron 6"
FT misc_feature 4076..4088
FT /tag= u
FT /note= "indefinite interval within intron 6"
FT exon 4225..4364
FT /tag= v
FT /note= "exon 7"
FT intron 4365..6571
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FT /note= "intron 7"
FT misc_feature 4391..4392
FT /tag= x
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FT /note= "known polymorphic site"
FT exon 6572..6677
FT /tag= aa
FT /note= "exon 8"
FT intron 6678..9163
FT /tag= ab
FT /note= "intron 8"
FT misc_feature 6823
FT /tag= ac
FT /note= "known polymorphic site"
FT misc_feature 9106
FT /tag= ad
FT /note= "known polymorphic site"
FT exon 9164..9209
FT /tag= ae
FT /note= "exon 9"
FT misc_feature 9207
FT /tag= af
FT /note= "known polymorphic site"
FT intron 9210..10530
FT /tag= ag
FT /note= "intron 9"
FT misc_feature 9376
FT /tag= ah
FT /note= "known polymorphic site"
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FT /tag= ai
FT /note= "exon 10"
FT 10608..11597
FT /tag= aj
FT /note= "intron 10"
FT 11384..11396
FT /tag= ak
FT /note= "indefinite interval within intron 10"
FT 11598..15023
FT /tag= al
FT /note= "exon 11"
FT 11908
FT /tag= am
FT /note= "known polymorphic site"
FT 11994
FT /tag= an
FT /note= "known polymorphic site"
FT 12952
FT /tag= ao
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FT 14041
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FT 14046
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FT /note= "known polymorphic site"
FT 14475
FT /tag= ay
FT /note= "known polymorphic site"
FT 14874
FT /tag= az
FT /note= "known polymorphic site"
FT 14891
FT /tag= ba
FT /note= "known polymorphic site"
FT 14966
FT /tag= bb
FT /note= "known polymorphic site"
FT 15024..15424
FT /tag= bc
FT /note= "intron 11"
FT 15284
FT /tag= bd
FT /note= "known polymorphic site"
FT 15425..15511
FT /tag= be
FT /note= "exon 12"
FT 15512..15952
FT /tag= bf
FT /note= "intron 12"
FT 15647..15659
FT /tag= bg
FT /note= "indefinite interval within intron 12"
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FT /tag= bh
FT /note= "exon 13"
FT 16077
FT /tag= bi
FT /note= "known polymorphic site"
FT 16127..16565
FT /tag= bj
FT /note= "intron 13"
FT 16243
FT /tag= bk
FT /note= "known polymorphic site"
FT 16370..16382
FT /tag= bl
FT /note= "known polymorphic site"

alignment_scores:
  Quality: 37.00 Length: 11
  Ratio: 3.700 Gaps: 0
  Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:
  US-08-653-294-18 x TL7519/rev ..
  Align seg 1/1 to reverse of: TL7519 from: 1 to: 24026
  2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
  |||||
  11314 AGATTGCCATCAGAAACTGGTCTCGATGTAC 11282

seq_name: N_Geneseq_36:TL7521

seq_documentation_block:
  ID TL7521 standard; cDNA; 24026 BP.
  AC TL7521;
  DT 04-OCR-1996 (first entry)
  DE Mutated BRCA1 genomic sequence from PMA08.1.
  KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
  KW antibody production; germline alteration; probe; lesion neoplasia; human;
  KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
  OS Homo sapiens.
  FH Key Location/Qualifiers
  FT exon 256..355
  FT /tag= a
  FT /note= "exon 1"
  FT 356..1512
  FT /tag= b
  FT /note= "intron 1"
  FT 1295
  FT /tag= c
  FT /note= "known polymorphic site"
  FT 1513..1611
  FT /tag= d
  FT /note= "exon 2"
  FT 1612..2206
  FT /tag= e
  FT /note= "intron 2"
  FT 1925..1937
  FT /tag= f
  FT /note= "indefinite interval within intron 2"
  FT 2141
  FT /tag= g
  FT /note= "known polymorphic site"
  FT 2207..2260
  FT /tag= h
  FT /note= "exon 3"
  FT 2261..2677
  FT /tag= i
  FT /note= "intron 3"
  FT 2569..2581
  FT /tag= j
  FT /note= "indefinite interval within intron 3"
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FT 2789. 3328
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FT 3329. 3406
FT /*tag= o
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FT 3407. 3813
FT /*tag= p
FT /note= "intron 5"
FT 3598. 3610
FT /*tag= q
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FT 3653
FT /*tag= r
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FT 3814. 3902
FT /*tag= s
FT /note= "exon 6"
FT 3903. 4224
FT /*tag= t
FT /note= "intron 6"
FT 4076. 4088
FT /*tag= u
FT /note= "indefinite interval within intron 6"
FT 4225. 4364
FT /*tag= v
FT /note= "exon 7"
FT 4385. 6571
FT /*tag= w
FT /note= "intron 7"
FT 4391. 4392
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FT 4602. 4614
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FT /note= "indefinite interval within intron 7"
FT 6538
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FT /note= "C to T mutation at known polymorphic site"
FT 6572. 6677
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FT /note= "exon 8"
FT 6678. 9163
FT /*tag= ab
FT /note= "intron 8"
FT 6823
FT /*tag= ac
FT /note= "known polymorphic site"
FT 9106
FT /*tag= ad
FT /note= "known polymorphic site"
FT 9164. 9209
FT /*tag= ae
FT /note= "exon 9"
FT 9207
FT /*tag= af
FT /note= "known polymorphic site"
FT 9210. 10530
FT /*tag= ag
FT /note= "intron 9"
FT 9376
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FT /*tag= ai
FT /note= "exon 10"
FT 10608. 11597
FT /*tag= aj
FT /note= "intron 10"
FT 11384. 11396
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FT /note= "indefinite interval within intron 10"
FT 11598. 115023
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FT 11908
FT /*tag= am
FT /note= "known polymorphic site"
FT 11994
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FT 14966
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FT 15024. 15424
FT /*tag= bc
FT /note= "intron 11"
FT 15284
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FT 15425. 15511
FT /*tag= be
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FT 15512. 15952
FT /*tag= bf
FT /note= "intron 12"
FT 15647. 15659
FT /*tag= bg
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FT exon /note= "indefinite interval within intron 12"
FT 15953. .16126
FT /tag= bh
FT /note= "exon 13"
FT misc_feature 16077
FT /tag= bi
FT /note= "known polymorphic site"
FT intron 16127. .16565
FT /tag= bj
FT /note= "intron 13"
FT misc_feature 16243
FT /tag= bk
FT /note= "known polymorphic site"
FT misc_feature 16370. .16382
FT /tag= bl

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alignment_scores: Quality: 37.00 Length: 11
 Ratio: 3.700 Gaps: 0
 Percent Similarity: 90.909 Percent Identity: 63.636

alignment_block:

US-08-653-294-18 x T17521/rev ..

Align seg 1/1 to reverse of: T17521 from: 1 to: 24026

2 ArgLeuAlaileArgArgileLeuLeuArgTyr 12

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|||||
11314 AGATTGGCCATCAGAAACGTGTTCTGATGATC 11282

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seq_name: N_Geneseq_36:T17522

seq_documentation_block:

ID T17522 standard; cDNA; 24026 BP.

AC T17522;

DT 04-OCT-1996 (first entry)

DE Mutated BRCA1 genomic sequence from PMA08.2.
 KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
 KW antibody production; germline alteration; probe; lesion neoplasia; human;
 KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
 OS Homo sapiens..

FH Key Location/Qualifiers

FT exon 256. .355

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FT /note= "exon 1"

FT intron 356. .1512

FT /tag= b

FT /note= "intron 1"

FT misc_feature 1295

FT /tag= c

FT /note= "known polymorphic site"

FT exon 1513. .1611

FT /tag= d

FT /note= "exon 2"

FT intron 1612. .2206

FT /tag= e

FT /note= "intron 2"

FT misc_feature 1925. .1937

FT /tag= f

FT /note= "indefinite interval within intron 2"

FT misc_feature 2141

FT /tag= g

FT /note= "known polymorphic site"

FT exon 2207. .2260

FT /tag= h

FT /note= "exon 3"

FT intron 2261. .2677

FT /tag= i

FT /note= "intron 3"

FT misc_feature 2569. .2581

FT /tag= j

FT /note= "indefinite interval within intron 3"

FT exon 2678. .2788

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FT /tag= k
FT /note= "exon 4"
FT 2725
FT /tag= l
FT /note= "known polymorphic site"
FT intron 2789. .3328
FT /tag= m
FT /note= "intron 4"
FT misc_feature 3063. .3075
FT /tag= n
FT /note= "indefinite interval within intron 4"
FT exon 3329. .3406
FT /tag= o
FT /note= "exon 5"
FT intron 3407. .3813
FT /tag= p
FT /note= "intron 5"
FT misc_feature 3598. .3610
FT /tag= q
FT /note= "indefinite interval within intron 5"
FT misc_feature 3653
FT /tag= r
FT /note= "known polymorphic site"
FT exon 3814. .3902
FT /tag= s
FT /note= "exon 6"
FT intron 3903. .4224
FT /tag= t
FT /note= "intron 6"
FT misc_feature 4076. .4088
FT /tag= u
FT /note= "indefinite interval within intron 6"
FT exon 4225. .4364
FT /tag= v
FT /note= "exon 7"
FT intron 4365. .6571
FT /tag= w
FT /note= "intron 7"
FT misc_feature 4391. .4392
FT /tag= x
FT /note= "known polymorphic site"
FT misc_feature 4602. .4614
FT /tag= y
FT /note= "indefinite interval within intron 7"
FT misc_feature 6538
FT /tag= z
FT /note= "known polymorphic site"
FT exon 6572. .6677
FT /tag= aa
FT /note= "exon 8"
FT intron 6678. .9163
FT /tag= ab
FT /note= "intron 8"
FT mutation 6823
FT /tag= ac
FT /note= "A to T mutation at known polymorphic site"
FT misc_feature 9106
FT /tag= ad
FT /note= "known polymorphic site"
FT exon 9164. .9209
FT /tag= ae
FT /note= "exon 9"
FT misc_feature 9207
FT /tag= af
FT /note= "known polymorphic site"
FT intron 9210. .10530
FT /tag= ag
FT /note= "intron 9"
FT misc_feature 9376
FT /tag= ah
FT /note= "known polymorphic site"
FT exon 10531. .10607
FT /tag= ai

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FT 10608..11597 /tag= aj
FT /note= "intron 10"
FT 11384..11396 /tag= ak
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FT 11598..15023 /tag= al
FT /note= "exon 11"
FT 11908 /tag= am
FT /note= "known polymorphic site"
FT 11994 /tag= an
FT /note= "known polymorphic site"
FT 12952 /tag= ao
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FT 15024..15424 /tag= bc
FT /note= "intron 11"
FT 15284 /tag= bd
FT /note= "known polymorphic site"
FT 15425..15511 /tag= be
FT /note= "exon 12"
FT 15512..15952 /tag= bf
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FT 15647..15659 /tag= bg
FT /note= "indefinite interval within intron 12"
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FT exon 15953..16126 /tag= bh
FT /note= "exon 13"
FT 16077 /tag= bi
FT /note= "known polymorphic site"
FT 16127..16565 /tag= bj
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FT 16243 /tag= bk
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FT 16370..16382 /tag= bl
FT alignment_scores:
Quality: 37.00 Length: 11
Ratio: 3.700 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636
alignment_block:
US-08-653-294-18 x TI17522/rev ..
Align seg 1/1 to reverse of: TI17522 from: 1 to: 24026
2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
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11314 AGATTGCCATCAGAAACCTGTTCTGATGTAC 11282
seq_name: N_Geneseq_36:TI17523
seq_documentation_block:
ID TI17523 standard; cDNA; 24026 BP.
AC TI17523;
DT 04-OCT-1996 (first entry)
DE Mutated BRCA1 genomic sequence from PMA09.2
KW Cancer therapy; breast and ovarian cancer predisposing gene; immunogen;
KW antibody production; germline alteration; probe; lesion neoplasia; human;
KW gene therapy; protein replacement therapy; protein mimetic; BRCA1; ds.
OS Homo sapiens.
FH Key Location/Qualifiers
FT exon 256..355 /tag= a
FT /note= "exon 1"
FT 356..1512 /tag= b
FT /note= "intron 1"
FT 1295 /tag= c
FT /note= "known polymorphic site"
FT 1513..1611 /tag= d
FT /note= "exon 2"
FT 1612..2206 /tag= e
FT /note= "intron 2"
FT 1925..1937 /tag= f
FT /note= "indefinite interval within intron 2"
FT 2141 /tag= g
FT /note= "known polymorphic site"
FT 2207..2260 /tag= h
FT /note= "exon 3"
FT 2261..2677 /tag= i
FT /note= "intron 3"
FT 2569..2581 /tag= j
FT /note= "indefinite interval within intron 3"
FT 2678..2788 /tag= k
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FT 2725 /tag= l
FT /note= "known polymorphic site"
FT intron 2789..3328
FT /tag= m
FT /note= "intron 4"
FT misc_feature 3063..3075
FT /tag= n
FT /note= "indefinite interval within intron 4"
FT exon 3329..3406
FT /tag= o
FT /note= "exon 5"
FT intron 3407..3813
FT /tag= p
FT /note= "intron 5"
FT misc_feature 3598..3610
FT /tag= q
FT /note= "indefinite interval within intron 5"
FT misc_feature 3653..
FT /tag= r
FT /note= "known polymorphic site"
FT exon 3814..3902
FT /tag= s
FT /note= "exon 6"
FT intron 3903..4224
FT /tag= t
FT /note= "intron 6"
FT misc_feature 4076..4088
FT /tag= u
FT /note= "indefinite interval within intron 6"
FT exon 4225..4364
FT /tag= v
FT /note= "exon 7"
FT intron 4365..6571
FT /tag= w
FT /note= "intron 7"
FT misc_feature 4391..4392
FT /tag= x
FT /note= "known polymorphic site"
FT misc_feature 4602..4614
FT /tag= y
FT /note= "indefinite interval within intron 7"
FT misc_feature 6538..
FT /tag= z
FT /note= "known polymorphic site"
FT exon 6572..6677
FT /tag= aa
FT /note= "exon 8"
FT intron 6678..9163
FT /tag= ab
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FT misc_feature 6823..
FT /tag= ac
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FT exon 9164..9209
FT /tag= ae
FT /note= "exon 9"
FT misc_feature 9207..
FT /tag= af
FT /note= "known polymorphic site"
FT intron 9210..10530
FT /tag= ag
FT /note= "intron 9"
FT misc_feature 9376..
FT /tag= ah
FT /note= "known polymorphic site"
FT mutation 9376..
FT /tag= ah
FT /note= "t to C mutation at known polymorphic site"
FT 10531..10607
FT /tag= ai
FT /note= "exon 10"
FT intron 10608..11597
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FT misc_feature 11384..11396
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FT exon 11598..15023
FT /tag= al
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FT misc_feature 11908..
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FT misc_feature 11994..
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FT misc_feature 14874..
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FT misc_feature 14891..
FT /tag= ba
FT /note= "known polymorphic site"
FT misc_feature 14966..
FT /tag= bb
FT /note= "known polymorphic site"
FT intron 15024..15424
FT /tag= bc
FT /note= "intron 11"
FT misc_feature 15284..
FT /tag= bd
FT /note= "known polymorphic site"
FT exon 15425..15511
FT /tag= be
FT /note= "exon 12"
FT intron 15512..15952
FT /tag= bf
FT /note= "intron 12"
FT misc_feature 15647..15659
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FT      /*tag= bg
FT      /note= "indefinite interval within intron 12"
FT      exon      15953..16126
FT      /*tag= bh
FT      /*tag= "exon 13"
FT      misc_feature 16077
FT      /*tag= bi
FT      /note= "known polymorphic site"
FT      intron      16127..16565
FT      /*tag= bj
FT      /note= "intron 13"
FT      misc_feature 16243
FT      /*tag= bk

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alignment_scores:
  Quality: 37.00      Length: 11
  Ratio: 3.700      Gaps: 0
Percent Similarity: 90.909 Percent Identity: 63.636

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alignment_block:
US-08-653-294-18 x T17523/rev ..
Align seg 1/1 to reverse of: T17523 from: 1 to: 24026

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```

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||
11314 AGATTGGCCATCAGAAACTGGTCTGATGTAC 11282

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OM of: US-08-653-294-18 to: EST:* out_format : pfs

Date: Feb 8, 2000 4:03 AM

About: Results were produced by the GenCore software, version 4.5.
Copyright (c) 1993-2000 CompuGen Ltd.

Command line parameters:
-MODEL=frame+p2n.model -DEV=xlp
-O=/cgnl_1/USPTO.spool/US08653294/funat_04022000_15770/app_query.fasta.1
-DB=EST -QMT=fastap -SUFFIX=rst -CAPOP=12.000 -GAPEXT=4.000
-MINMATCH=0.100 -LOOCL=0.000 -LOOPEXT=0.000 -QGAPOP=4.500
-QGAPEXT=0.050 -XGAPOP=10.000 -YGAPEXT=0.500 -FGAPOP=6.000
-FGAPEXT=7.000 -START=1 -MATRIX=blossum62 -TRANS=human40.cdi
-DELEX=7.000 -ALIGN=200 -THR SCORE=pct -ALIGN=15 -MODE=LOCAL
-OUTPMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=1000000 -USER=US08653294
-NCPU=6 -ICPU=3 -NO_XLPXY -WAIT -THREADS=1

Search information block:

Query: US-08-653-294-18

Query length: 12

Database: EST:*

Database sequences: 4538634

Database length: 1887831982

Search time (sec): 8553.360000

score_list:

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gb_gss6:AQ869567	+	42.00	141.78	39.36	438	AQ869567 nbbe0035D05f CUGI Rice
gb_gss6:AQ862871	+	42.00	140.92	43.94	483	AQ862871 nbbe0019L13f CUGI Rice
gb_est34:AV120004	+	40.00	139.07	55.68	259	AV120004 AV120004 Mus musculus
gb_est36:AV182651	+	40.00	137.78	65.69	300	AV182651 AV182651 Yujii Kohara
gb_gss6:AQ864125	+	40.00	135.02	93.62	411	AQ864125 nbbe0022F09f CUGI Rice
gb_gss6:AQ864125	+	40.00	134.37	101.86	443	AQ864125 nbbe0004H21f CUGI Rice
gb_gss6:AQ864125	+	40.00	134.03	106.26	460	AQ864125 nbbe0005D22f CUGI Rice
gb_est24:AV1235399	+	40.00	133.27	117.24	502	AV1235399 EST231961 Normalized
gb_est27:AV1407553	+	40.00	133.11	119.61	511	AV1407553 EST235843 Normalized
gb_gss15:AQ861929	+	40.00	132.98	121.72	519	AQ861929 HS_2130_B2_H03_MR CIT
gb_gss7:AA917001	+	40.00	132.93	122.51	522	AA917001 nbbe0067L07f CUGI Rice
gb_gss6:AQ870759	+	40.00	129.17	198.33	801	AQ870759 nbbe0004H21f CUGI Rice
gb_gss13:AQ279730	+	39.00	133.90	108.16	308	AQ279730 CITB1-E1-2513B14.TR CIT
gb_est19:AA748739	+	39.00	133.87	108.55	309	AA748739 ny06e12.s1 NCI_CGAP GC
gb_est14:AA389896	+	39.00	132.50	129.31	361	AA389896 vb30f04.r1 Soares mous
gb_est44:AW214737	+	39.00	131.83	141.05	390	AW214737 uc99c07.y1 NCI_CGAP_Lu
gb_est9:AA119739	+	39.00	130.79	161.15	439	AA119739 mnl14d02.r1 Beddington
gb_est14:AA414705	+	39.00	130.77	161.56	440	AA414705 vc69c04.s1 Knowles Sol
gb_est18:AA692200	+	39.00	130.23	173.17	468	AA692200 vt19h09.r1 Barstead m
gb_gss6:AQ863628	+	39.00	130.17	174.42	471	AQ863628 nbbe0021F21f CUGI Rice
gb_est20:AA839058	+	39.00	129.90	180.68	486	AA839058 vw47a11.r1 Soares mous
gb_est8:AA012276	+	39.00	129.88	181.10	487	AA012276 TgESTz16c11.r1 TgME49
gb_gss13:AQ838867	+	39.00	129.79	183.20	492	AQ838867 HS_5077_B2_H02_T7A RQC
gb_est10:AA168371	+	39.00	129.77	183.62	493	AA168371 mr28a04.r1 Soares fetal
gb_est15:H72837	+	39.00	129.58	188.23	504	H72837 vs06c01.s1 Soares mous
gb_est12:AA289553	+	39.00	129.56	188.65	505	AA289553 vb17b05.r1 Soares mous
gb_est11:AA239865	+	39.00	129.52	189.49	507	AA239865 mx80f12.r1 Soares mous
gb_est9:AA103036	+	39.00	129.32	194.55	519	AA103036 mo2id11.r1 Life Tech m
gb_est36:AA1891896	+	39.00	128.29	195.39	521	AA1891896 ul60c04.x1 Sugano mous
gb_est10:AA174998	+	39.00	128.21	224.31	589	AA174998 ms88h04.r1 Soares mous
gb_est20:AA869826	+	39.00	128.21	224.31	589	AA869826 vq16c07.r1 Barstead st
gb_gss14:AA569045	+	39.00	128.11	227.31	596	AA569045 HS_5344_A1_F08_T7A RQC
gb_est10:AA137602	+	39.00	127.97	231.17	605	AA137602 mq9a02.r1 Soares mous
gb_est17:AA638393	+	39.00	127.72	238.92	623	AA638393 vl99d11.r1 Knowles Sol
gb_est16:AA571878	+	39.00	127.55	244.11	635	AA571878 vm04e12.r1 Knowles Sol
gb_gss15:AQ856284	+	39.00	127.54	244.54	636	AQ856284 Sheared DNA-27K23.TF4S
gb_gss15:AQ856284	+	39.00	127.51	254.95	660	AQ856284 Sheared DNA-5K21.TF SH
gb_gss15:AQ856284	+	39.00	127.51	254.95	660	AQ856284 Sheared DNA-5K21.TF SH
gb_gss13:R08474	+	39.00	122.50	466.40	1129	R08474 TL2N24-T7 TAMU Arabidop
gb_est10:AA171093	+	38.00	137.41	68.92	136	AA171093 ms50c07.r1 Life Tech m
gb_est39:AA124405	+	38.00	133.52	113.56	212	AA124405 UI-M-BH2.1-ape-d-08-0
gb_est34:AV135435	+	38.00	130.53	166.57	298	AV135435 AV135435 Mus musculus

gb_est33:AV117866 - 38.00 130.33 170.98 305 ! AV117866 AV117866 Mus muscul
gb_est23:AU022889 - 38.00 128.68 211.20 368 ! AU022889 AU022889 Mouse unfe
gb_gss12:AQ366741 + 38.00 127.60 242.44 416 ! AQ366741 HS_5038_B2_D09_SP6E

seq_name: gb_gss9:AQ137850

seq_documentation_block:
LOCUS AQ137850 461 bp DNA GSS 24-SEP-1998
DEFINITION HS_3058_B2_C09_MF CIT Approved Human Genomic Sperm Library D Homo
sapiens genomic clone plate=3058 Col=18 Row=F, genomic survey
sequence.
ACCESSION AQ137850
VERSION AQ137850.1 GI:3528503
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 461)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589
COMMENT Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3058 row: F column: 18
Class: BAC ends
High quality sequence stop: 461.
Location/Qualifiers
1. 461
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone_lib="CIT Approved Human Genomic Sperm Library D"
/clone="Plate:3058 Col=18 Row=F"
/sex="male"
/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coli DH10B"

BASE COUNT 132 a 125 c 102 g 102 t
ORIGIN

alignment_scores:
Quality: 44.00 Length: 12
Ratio: 4.000 Gaps: 0
Percent Similarity: 91.667 Percent Identity: 75.000

alignment_block:
US-08-653-294-18 x AQ137850 ..
Align seg 1/1 to: AQ137850 from: 1 to: 461

1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
:::|||||:::|||||:::|||||:::|||||
18 CATCGACTCACTAAGGCGAATCTCGCCAGGTAC 53

seq_name: gb_gss6:AQ869567

seq_documentation_block:
LOCUS AQ869567 438 bp DNA GSS 03-NOV-1999
DEFINITION nbbe0035D05f CUGI Rice BAC Library (EcoRI) Oryza sativa genomic
clone nbbe0035D05f, genomic survey sequence.
ACCESSION AQ869567
VERSION AQ869567.1 GI:6220018
KEYWORDS GSS.

SOURCE
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.
REFERENCE 1 (bases 1 to 438)
AUTHORS Wing,R.A. and Dean,R.A.
TITLE A BAC End Sequencing Framework to Sequence the Rice Genome
JOURNAL Unpublished (1998)
COMMENT On Mar 23, 1999 this sequence version replaced gi:3324665.
Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Seq primer: TAATACGACTCACTATAGGG
Class: BAC ends
High quality sequence start: 24
High quality sequence stop: 397.
FEATURES
source
1..438
/organism="Oryza sativa"
/strain="Japonica"
/cultivar="Nipponbare"
/db_xref="taxon:4530"
/clone="nbeb0035D05f"
/clone_lib="CUGI Rice BAC Library (EcoRI)"
/tissue_type="Leaf"
/lab_host="E. coli DH10B"
/note="vector: pBACindigo; Site_1: EcoRI; Site_2: EcoRI; Rice is the most important food crop in the world. Half of the world population, especially those inhabiting highly populated areas of the humid tropics and subtropics, rely on rice as their primary source of carbohydrate. Monocotyledonous rice is a diploid plant (2n=24) with a haploid genome equivalent of 431 Mbp (Arumuganathan and Earle, 1991). The relatively small genome of rice, three times larger than that of Arabidopsis, makes it suitable for genomic studies. In order to facilitate positional cloning, physical mapping and genome sequencing of rice, we have constructed a BAC library from Oryza sativa. Nipponbare variety using EcoRI as the cloning enzyme. The library contains 55,296 clones with an average insert size of 121 kb providing approximately 15 haploid genome equivalents. The deep coverage allows the isolation a particular sequence with a probability of 99.9 %. Three high density filters, each containing 18,432 clones (doubly spotted), represent the whole library for colony screening and can be requested from the Clemson University BAC/EST Resource Center (www.genome.clemson.edu)."

BASE COUNT 121 a 86 c 65 g 164 t 2 others
ORIGIN

alignment_scores:
Quality: 42.00 Length: 10
Ratio: 4.200 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 90.000

alignment_block:
US-08-653-294-18 x AQ869567 ..
Align seg 1/1 to: AQ869567 from: 1 to: 438

3 LeuAlaIleArgArgIleLeuLeuArgTyr 12
||||:|||||
15 CTCATCATACGCGGCAATCTATTAGATAT 44

seq_name: gb_gss6:AQ862871
seq_documentation_block:

LOCUS
DEFINITION AQ862871 483 bp DNA GSS 03-NOV-1999
clone nbeb0019L13f CUGI Rice BAC Library (EcoRI) Oryza sativa genomic
ACCESSION AQ862871
VERSION
KEYWORDS AQ862871.1 GI:6213328
SOURCE
ORGANISM Oryza sativa.
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.
REFERENCE 1 (bases 1 to 483)
AUTHORS Wing,R.A. and Dean,R.A.
TITLE A BAC End Sequencing Framework to Sequence the Rice Genome
JOURNAL Unpublished (1998)
COMMENT Contact: Wing RA
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Seq primer: TAATACGACTCACTATAGGG
Class: BAC ends
High quality sequence start: 40
High quality sequence stop: 411.
FEATURES
source
1..483
/organism="Oryza sativa"
/strain="Japonica"
/cultivar="Nipponbare"
/db_xref="taxon:4530"
/clone="nbeb0019L13f"
/clone_lib="CUGI Rice BAC Library (EcoRI)"
/tissue_type="Leaf"
/lab_host="E. coli DH10B"
/note="vector: pBACindigo; Site_1: EcoRI; Site_2: EcoRI; Rice is the most important food crop in the world. Half of the world population, especially those inhabiting highly populated areas of the humid tropics and subtropics, rely on rice as their primary source of carbohydrate. Monocotyledonous rice is a diploid plant (2n=24) with a haploid genome equivalent of 431 Mbp (Arumuganathan and Earle, 1991). The relatively small genome of rice, three times larger than that of Arabidopsis, makes it suitable for genomic studies. In order to facilitate positional cloning, physical mapping and genome sequencing of rice, we have constructed a BAC library from Oryza sativa. Nipponbare variety using EcoRI as the cloning enzyme. The library contains 55,296 clones with an average insert size of 121 kb providing approximately 15 haploid genome equivalents. The deep coverage allows the isolation a particular sequence with a probability of 99.9 %. Three high density filters, each containing 18,432 clones (doubly spotted), represent the whole library for colony screening and can be requested from the Clemson University BAC/EST Resource Center (www.genome.clemson.edu)."

BASE COUNT 132 a 96 c 101 g 151 t 3 others
ORIGIN

alignment_scores:
Quality: 42.00 Length: 11
Ratio: 4.200 Gaps: 0
Percent Similarity: 90.909 Percent Identity: 81.818

alignment_block:
US-08-653-294-18 x AQ862871 ..
Align seg 1/1 to: AQ862871 from: 1 to: 483

2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
||||:|||||
||||:|||||

22 AGACTCTCTATACGGCGAATCTTTAGCCTAT 54

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seq_name: gb_est34:AV120004

seq_documentation_block:
  LOCUS      AV120004      259 bp      mRNA      EST      30-JUN-1999
  DEFINITION Mus musculus C57BL/6J 10-day embryo Mus musculus cDNA
  clone 2610307A11, mRNA sequence.
  ACCESSION  AV120004
  VERSION     AV120004.1 GI:5302155
  KEYWORDS   EST.
  SOURCE     house mouse.
  ORGANISM   Mus musculus

REFERENCE
  1 (bases 1 to 259)
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
  Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS
  Carninci, P., Shibata, K., Ozawa, Y., Konno, H., Itoh, M., Aizawa, K.,
  Akahira, S., Akiyama, J., Fukuda, S., Fukunishi, Y., Funayama, T.,
  Hara, A., Hayatsu, N., Hori, F., Ishikawa, T., Itoh, M., Izawa, M.,
  Kawai, J., Kikuchi, N., Kojima, Y., Matsuyama, T., Niitsuma, H., Oda, H.,
  Owa, C., Sato, K., Shibata, Y., Shigemoto, Y., Shiraki, T., Sogabe, Y.,
  Sugahara, Y., Suzuki, H., Suzuki, H., Tateno, M., Tomaru, Y.,
  Tomimaga, N., Watanabe, S., Yagame, M., Yamamura, T., Yokota, T.,
  Yoshino, M., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.

TITLE
  RIKEN Mouse ESTs
JOURNAL
  Unpublished (1999)
COMMENT
  On Jun 5, 1998 this sequence version replaced gi:3188908.
  Contact: Chie Owa
  Genome Science Laboratory
  RIKEN
  3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan
  Tel: 81-298-36-9145
  Fax: 81-298-36-9098
  Email: genome-resetc.riken.go.jp
  Thermostabilization and thermoactivation of thermolabile enzymes by
  trehalose and its application for the synthesis of full length cDNA
  (Proc. Natl. Acad. Sci. U.S.A. 95(2):520-524 (1998))
  Transcriptional sequencing: A method for DNA sequencing using RNA
  polymerase (Proc. Natl. Acad. Sci. U.S.A. 95(7):3455-3460 (1998))
  Please visit our web site (http://genome.rtc.riken.go.jp) for
  further details.

FEATURES
  source
    location/Qualifiers
      1..259
        /organism="Mus musculus"
        /strain="C57BL/6J"
        /db_xref="taxon:10090"
        /clone="2610307A11"
        /clone_lib="Mus musculus C57BL/6J 10-day embryo"
        /sex="mixed"
        /dev_stage="10-day embryo"
      70 a 57 c 48 g 84 t

BASE COUNT
  70 a 57 c 48 g 84 t
ORIGIN

alignment_scores:
  Quality: 40.00      Length: 11
  Ratio: 3.636       Gaps: 0
  Percent Similarity: 100.000 Percent Identity: 63.636

alignment_block:
  US-08-653-294-18 x AV120004/rev ..
  Align seg 1/1 to reverse of: AV120004 from: 1 to: 259

  2 ArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
  ::::::::::::::::::::::::::::::::::::
  241 AGATTAGCCCTTAAAGAGTCTTATAAATAT 209

seq_name: gb_est36:AV182651
seq_documentation_block:
  LOCUS      AV182651      300 bp      mRNA      EST      21-JUL-1999
  DEFINITION AV182651 Yuji Kohara unpublished cDNA:Strain N2 hermaphrodite
  clone 2610307A11, mRNA sequence.
  ACCESSION  AV182651
  VERSION     AV182651.1 GI:5302155
  KEYWORDS   EST.
  SOURCE     house mouse.
  ORGANISM   Mus musculus

REFERENCE
  1 (bases 1 to 300)
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
  Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

AUTHORS
  Carninci, P., Shibata, K., Ozawa, Y., Konno, H., Itoh, M., Aizawa, K.,
  Akahira, S., Akiyama, J., Fukuda, S., Fukunishi, Y., Funayama, T.,
  Hara, A., Hayatsu, N., Hori, F., Ishikawa, T., Itoh, M., Izawa, M.,
  Kawai, J., Kikuchi, N., Kojima, Y., Matsuyama, T., Niitsuma, H., Oda, H.,
  Owa, C., Sato, K., Shibata, Y., Shigemoto, Y., Shiraki, T., Sogabe, Y.,
  Sugahara, Y., Suzuki, H., Suzuki, H., Tateno, M., Tomaru, Y.,
  Tomimaga, N., Watanabe, S., Yagame, M., Yamamura, T., Yokota, T.,
  Yoshino, M., Muramatsu, M., Okazaki, Y. and Hayashizaki, Y.

TITLE
  RIKEN Mouse ESTs
JOURNAL
  Unpublished (1999)
COMMENT
  On Jun 5, 1998 this sequence version replaced gi:3188908.
  Contact: Chie Owa
  Genome Science Laboratory
  RIKEN
  3-1-1 Koyadai, Tsukuba, Ibaraki 305-0074, Japan
  Tel: 81-298-36-9145
  Fax: 81-298-36-9098
  Email: genome-resetc.riken.go.jp
  Thermostabilization and thermoactivation of thermolabile enzymes by
  trehalose and its application for the synthesis of full length cDNA
  (Proc. Natl. Acad. Sci. U.S.A. 95(2):520-524 (1998))
  Transcriptional sequencing: A method for DNA sequencing using RNA
  polymerase (Proc. Natl. Acad. Sci. U.S.A. 95(7):3455-3460 (1998))
  Please visit our web site (http://genome.rtc.riken.go.jp) for
  further details.

FEATURES
  source
    location/Qualifiers
      1..300
        /organism="Caenorhabditis elegans"
        /strain="N2"
        /db_xref="taxon:6239"
        /clone="yk640g7"
        /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
        hermaphrodite embryo"
        /sex="hermaphrodite"
        /dev_stage="embryo"
      105 a 54 g 71 t 2 others

BASE COUNT
  105 a 54 g 71 t 2 others
ORIGIN

seq_name: gb_gss6:AQ864125
seq_documentation_block:
  LOCUS      AQ864125      411 bp      DNA      GSS      03-NOV-1999
  DEFINITION nbe0022F09f CUGI Rice BAC Library (EcoRI) Oryza sativa genomic
  clone nbe0022F09f, genomic survey sequence.
  ACCESSION  AQ864125
  VERSION     AQ864125.1 GI:6214582
  KEYWORDS   GSS.
  SOURCE     Oryza sativa.
  ORGANISM   Oryza sativa
  Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
  euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
  Poaceae; Oryza.
  1 (bases 1 to 411)
  Wing, R.A. and Dean, R.A.
  A BAC End Sequencing Framework to Sequence the Rice Genome
  Unpublished (1998)
  Contact: Wing RA
  Clemson University Genomics Institute
  Clemson University
  100 Jordan Hall, Clemson, SC 29634, USA

```

```

embryo Caenorhabditis elegans cDNA clone yk640g7 3', mRNA sequence.
AV182651
AV182651.1 GI:5562552
EST.
Caenorhabditis elegans.
Caenorhabditis elegans
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 300)
Kohara, Y., Shin-i, T., Thierry-Mieg, J., Thierry-Mieg, D., Mitsuki, H.,
Nishigaki, A., Motoshita, T., Zeng, Q., Watanabe, H., Sugimoto, A., and
Sano, M., Miyata, A., Mitani, Y., Iida, K., Uesugi, H., Sugiyama, Y. and
Nomoto, H.
Expressed genes in C.elegans
Unpublished (1999)
On Jun 5, 1998 this sequence version replaced gi:3189500.
Contact: Yuji Kohara
Gene Library Lab
National Institute of Genetics
Yata 1111, Mishima, Shizuoka 411, Japan
Tel: 81-559-81-6854
Fax: 81-559-81-6855
Email: ykohara@lab.nig.ac.jp.
Location/Qualifiers
  1..300
    /organism="Caenorhabditis elegans"
    /strain="N2"
    /db_xref="taxon:6239"
    /clone="yk640g7"
    /clone_lib="Yuji Kohara unpublished cDNA:Strain N2
    hermaphrodite embryo"
    /sex="hermaphrodite"
    /dev_stage="embryo"
  105 a 54 g 71 t 2 others

BASE COUNT
  105 a 54 g 71 t 2 others
ORIGIN

alignment_scores:
  Quality: 40.00      Length: 12
  Ratio: 4.000       Gaps: 0
  Percent Similarity: 83.333 Percent Identity: 58.333

alignment_block:
  US-08-653-294-18 x AV182651 ..
  Align seg 1/1 to: AV182651 from: 1 to: 300

  1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
  ::::::::::::::::::::::::::::::::::::
  85 TACCGACTACTATCGGTAGTATTATAACAATAT 120

seq_name: gb_gss6:AQ864125
seq_documentation_block:
  LOCUS      AQ864125      411 bp      DNA      GSS      03-NOV-1999
  DEFINITION nbe0022F09f CUGI Rice BAC Library (EcoRI) Oryza sativa genomic
  clone nbe0022F09f, genomic survey sequence.
  ACCESSION  AQ864125
  VERSION     AQ864125.1 GI:6214582
  KEYWORDS   GSS.
  SOURCE     Oryza sativa.
  ORGANISM   Oryza sativa
  Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
  euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
  Poaceae; Oryza.
  1 (bases 1 to 411)
  Wing, R.A. and Dean, R.A.
  A BAC End Sequencing Framework to Sequence the Rice Genome
  Unpublished (1998)
  Contact: Wing RA
  Clemson University Genomics Institute
  Clemson University
  100 Jordan Hall, Clemson, SC 29634, USA

```

```

TITLE      Sequence-tagged connectors: A sequence approach to mapping and
           scanning the human genome
JOURNAL    Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE    99380589
COMMENT    Contact: Mahairas GG, Wallace JC, Hood L
           High Throughput Sequencing Center
           University of Washington
           401 Queen Anne Avenue North, Seattle, WA 98109, USA
           Tel: (206) 616-3618
           Fax: (206) 616-3887
           Email: jwallace@u.washington.edu
           Clones are derived from the human BAC library RPCI-11. For BAC
           library availability, please contact Pieter de Jong
           (pieter@dejong.med.buffalo.edu). Clones may be purchased from
           BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
           or from Resear h Genetics (info@resgen.com). BAC end Web Server:
           http://www.htsc.washington.edu
           Plate: 711 row: D column: 14
           Seq primer: SP6
           Class: BAC ends
           High quality sequence stop: 443.
FEATURES   Location/Qualifiers
            source
              1..443
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="Plate-711 Col-14 Row-D"
                /clone_lib="RPCI-11 Human Male BAC Library"
                /sex="male"
                /note="vector: pBACE3.6; Genomic sequence of BAC ends"
            BASE COUNT      134 a      95 c      83 g      130 t      1 others
ORIGIN
alignment_scores
  Quality: 40.00      Length: 12
  Ratio: 3.333      Gaps: 0
  Percent Similarity: 100.000      Percent Identity: 50.000
alignment_block:
  US-08-653-294-18 x AQ460667 ..
Align seg 1/1 to: AQ460667 from: 1 to: 443
1  TYZArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||:|||||:
237  TACCAATGACCTCAGAGACTTTTGTCGGGTTTC 272
seq_name: gb_gss6:AQ857430
seq_documentation_block:
LOCUS      AQ857430      460 bp      DNA      GSS      03-NOV-1999
DEFINITION nbe0005D22f CUGI Rice BAC Library (EcoRI) Oryza sativa genomic
           clone nbe0005D22f, genomic survey sequence.
ACCESSION  AQ857430
VERSION    AQ857430
KEYWORDS   GSS.
SOURCE     Oryza sativa.
ORGANISM   Oryza sativa
           Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
           euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
           Poaceae; Oryza.
1 (bases 1 to 460)
Wing, R.A. and Dean, R.A.
A BAC End Sequencing Framework to Sequence the Rice Genome
Unpublished (1998)
Contact: Wing RA
Clemson University
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Seq primer: TAAATCAGACTCATATAGGG

```


alignment_scores:
 Quality: 40.00 Length: 12
 Ratio: 4.444 Gaps: 0
 Percent Similarity: 75.000 Percent Identity: 75.000

alignment_block:

US-08-653-294-18 x AI407553/rev ..

Align seg 1/1 to reverse of: AI407553 from: 1 to: 511

1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
 |||||
 378 TACAGACTTGCTACAGCTTATCTATTAGGTAT 343

seq_name: gb_gss15:AQ661929

seq_documentation_block: 519 bp DNA GSS 23-JUN-1999
 LOCUS AQ661929 HS_2130_B2_H03_MR CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate-2130 Col-6 Row-P, genomic survey sequence.
 DEFINITION
 ACCESSION AQ661929
 VERSION AQ661929.1 GI:5169697
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 519)
 AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
 TITLE Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
 JOURNAL Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
 MEDLINE 99380589
 COMMENT On Sep 10, 1998 this sequence version replaced gi:3556096. Contact: Mahairas GG, Wallace JC, Hood L High Throughput Sequencing Center University of Washington 401 Queen Anne Avenue North, Seattle, WA 98109, USA Tel: (206) 616-3618 Fax: (206) 616-3887 Email: jwallace@u.washington.edu Clones may be purchased from Research Genetics (info@resgen.com). BAC end Web Server: http://www.htsc.washington.edu Plate: 2130 row: P column: 6 Seq primer: M13 Reverse Class: BAC ends High quality sequence stop: 519.

FEATURES

source
 1..519
 Location/Qualifiers
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="plate=2130 Col-6 Row-P"
 /clone_lib="CIT Approved Human Genomic Sperm Library D"
 /sex="male"
 /note="Organ: sperm; Vector: pBelOBAC11; BAC Clones in E-Coli DH10B"

BASE COUNT 169 a 106 c 108 g 136 t
 ORIGIN

alignment_scores:
 Quality: 40.00 Length: 10
 Ratio: 4.000 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-18 x AQ661929/rev ..

Align seg 1/1 to reverse of: AQ661929 from: 1 to: 519

1 TyrArgLeuAlaIleArgArgIleLeuLeu 10
 |||||
 154 TATCGTTTGTCTATTAGAGAGTGTGCTT 125

seq_name: gb_gss7:AQ917001

seq_documentation_block: 522 bp DNA GSS 02-DEC-1999
 LOCUS AQ917001 nbe0067L07f CUGI Rice BAC Library (ECORI) Oryza sativa genomic clone nbe0067L07f, genomic survey sequence.
 DEFINITION
 ACCESSION AQ917001
 VERSION AQ917001.1 GI:6513517
 KEYWORDS GSS.
 SOURCE Oryza sativa.

ORGANISM

Oryza sativa
 Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae; Oryza.

REFERENCE

1 (bases 1 to 522)

AUTHORS

Wing,R.A. and Dean,R.A.

TITLE

A BAC End Sequencing Framework to Sequence the Rice Genome

JOURNAL

Unpublished (1998)

COMMENT

Contact: Wing RA

Clemson University Genomics Institute

100 Jordan Hall, Clemson, SC 29634, USA

Tel: 864 656 7288

Fax: 864 656 4293

Email: rwing@clemson.edu

Seq primer: TAAACGACTCCTACTATAGGG

Class: BAC ends

High quality sequence start: 34

High quality sequence stop: 383.

FEATURES

source
 1..522
 Location/Qualifiers
 /organism="Oryza sativa"
 /strain="Japonica"
 /cultivar="Nipponbare"
 /db_xref="taxon:4530"
 /clone="nbe0067L07f"
 /clone_lib="CUGI Rice BAC Library (ECORI)"
 /tissue_type="Leaf"
 /lab_host="E. coli DH10B"

/note="Vector: pBACIndigo; Site:1: EcoRI; Site:2: EcoRI; Note is the most important food crop in the world. Half of the world population, especially those inhabiting highly populated areas of the humid tropics and subtropics, rely on rice as their primary source of carbohydrate. Monocotyledonous rice is a diploid plant (2n=24) with a haploid genome equivalent of 431 Mbp (Arumuganathan and Earle, 1991). The relatively small genome of rice, three times larger than that of Arabidopsis, makes it suitable for genomic studies. In order to facilitate positional cloning, physical mapping and genome sequencing of rice, we have constructed a BAC library from Oryza sativa, Nipponbare variety using EcoRI as the cloning enzyme. The library contains 55,296 clones with an average insert size of 121 kb providing approximately 15 haploid genome equivalents. The deep coverage allows the isolation a particular sequence with a probability of 99.9 %. Three high density filters, each containing 18,432 clones (doubly spotted), represent the whole library for colony screening and can be requested from the Clemson University BAC/EST Resource Center (www.genome.clemson.edu)."

BASE COUNT 154 a 103 c 101 g 163 t
 ORIGIN

alignment_scores:

Quality: 40.00 Length: 10
 Ratio: 4.000 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 90.000

alignment_block:

US-08-653-294-18 x AQ917001 ..

Align seg 1/1 to: AQ917001 from: 1 to: 522

2 ArgLeuAlaIleArgArgIleLeuArg 11

|||||:|||||:|||||:|||||:|||||

16 CGATTGACTATACGCCGAATTCCTACTACGA 45

seq_name: gb_gss6:AQ857059

seq_documentation_block:

LOCUS AQ857059 801 bp DNA GSS 03-NOV-1999
DEFINITION nbe0004H21f CUGI Rice BAC Library (EcoRI) Oryza sativa genomic
clone nbe0004H21f, genomic survey sequence.

ACCESSION AQ857059

VERSION AQ857059.1 GI:6207425

KEYWORDS GSS.

SOURCE Oryza sativa.

ORGANISM Oryza sativa

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
Poaceae; Oryza.

1 (bases 1 to 801)

Wing, R.A. and Dean, R.A.

A BAC End Sequencing Framework to Sequence the Rice Genome

JOURNAL

COMMENT

Contact: Wing RA

Clemson University Genomics Institute

Clemson University

100 Jordan Hall, Clemson, SC 29634, USA

Tel: 864 656 7288

Fax: 864 656 4293

Email: rwing@clemson.edu

Seq primer: TAATACGACTCTACTATAGG

Class: BAC ends

High quality sequence start: 24

High quality sequence stop: 360.

FEATURES

source

1..801

/organism="Oryza sativa"

/strain="Japonica"

/cultivar="Nipponbare"

/db_xref="taxon:4530"

/clone="nbe0004H21f"

/clone_lib="CUGI Rice BAC Library (EcoRI)"

/tissue_type="Leaf"

/lab_host="E. coli DH10B"

/note="Vector: pBACIndigo; Site_1: EcoRI; Site_2: EcoRI;
rice is the most important food crop in the world. Half of
the world population, especially those inhabiting highly
populated areas of the humid tropics and subtropics, rely
on rice as their primary source of carbohydrate.

Monocotyledonous rice is a diploid plant (2n=24) with a
haploid genome equivalent of 431 Mbp (Arumuganathan and
Earle, 1991). The relatively small genome of rice, three
times larger than that of Arabidopsis, makes it suitable
for genomic studies. In order to facilitate positional
cloning, physical mapping and genome sequencing of rice,
we have constructed a BAC library from Oryza sativa,
Nipponbare variety using EcoRI as the cloning enzyme. The
library contains 55,296 clones with an average insert size
of 121 kb providing approximately 15 haploid genome
equivalents. The deep coverage allows the isolation a
particular sequence with a probability of 99.9 %. Three
high density filters, each containing 18,432 clones
(doubly spotted), represent the whole library for colony
screening and can be requested from the Clemson University
BAC/EST Resource Center (www.genome.clemson.edu)."

BASE COUNT

ORIGIN

259 a 158 c 162 g 220 t

2 others

alignment_scores:

Quality: 40.00 Length: 10

Ratio: 4.000 Gaps: 0

Percent Similarity: 100.000 Percent Identity: 80.000

alignment_block:

US-08-653-294-18 x AQ857059 ..

Align seg 1/1 to: AQ857059 from: 1 to: 801

3 LeuAlaIleArgArgIleLeuArgTyr 12

|||||:|||||:|||||:|||||:|||||

15 CTCACATACGCGGAATTCCTATCGCTAC 44

seq_name: gb_gss11:AQ279730

seq_documentation_block:

LOCUS AQ279730 308 bp DNA GSS 22-NOV-1998
DEFINITION CITBI-EI-2513B14.TR CITBI-EI Homo sapiens genomic clone 2513B14,
genomic survey sequence.

ACCESSION AQ279730

VERSION AQ279730.1 GI:3905634

KEYWORDS GSS.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 308)

Adams, M.D., Rounsley, S.D., Zhao, S., Bass, S., Linher, K., Golden, K.,
Berry, K., Granger, D., Suh, E., Wibie, C., Shizuya, H., Simon, M. and
Venter, J.C.
Use of a random human BAC End Sequence Database for Sequence-Ready
Map Building

Unpublished (1998)

Other_GSSs: CITBI-EI-2513B14.TF

Contact: Mark Adams

Department of Eukaryotic Genomics

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850, USA

Tel: 301 838 0200

Fax: 301 838 0208

Email: mdadams@tigr.org

Clones are available from Research Genetics (info@resgen.com). BAC

end search page:

http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.

Seq primer: M13 Reverse

Class: BAC ends.

Location/Qualifiers

1..308

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="2513B14"

/clone_lib="CITBI-EI"

/sex="male"

/cell_type="sperm"

/note="Vector: pBelOBAC11; Site_1: EcoRI; Site_2: EcoRI;
Caltech Human BAC Library D"

75 a 76 c 55 g 102 t

BASE COUNT

ORIGIN

39.00 Length: 12

Ratio: 3.900 Gaps: 0

Percent Similarity: 83.333 Percent Identity: 58.333

alignment_scores:

Quality: 39.00 Length: 12

Ratio: 3.900 Gaps: 0

Percent Similarity: 83.333 Percent Identity: 58.333

alignment_block:

US-08-653-294-18 x AQ279730 ..

Align seg 1/1 to: AQ279730 from: 1 to: 308

1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
||| |||||::: ||::|||:::|||||
223 TACTCTCGCTTGTACAGACTGCTAATGAGATAC 258

seq_name: gb_est19:AA748739

seq_documentation_block:
LOCUS AA748739 309 bp mRNA EST 22-JAN-1998
DEFINITION nv06e12.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1270990 3',
mRNA sequence.

ACCESSION AA748739

VERSION AA748739.1 GI:2788697

KEYWORDS EST.

SOURCE

ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 309)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

On Jan 14, 1998 this sequence version replaced gi:1797612.

Contact: Robert Strausberg, Ph.D.

Tel.: (301) 496-1550

Email: Robert_Strausberg@nih.gov

Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,

Ph.D., Gerald Marti, M.D.

CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima

Bonaldo, Ph.D.

CDNA Library Arrayed by: Greg Lennon, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be

found through the I.M.A.G.E. Consortium/LLNL at:

www-bio.llnl.gov/bbrp/image/image.html

Insert Length: 574 Std Error: 0.00

Seq primer: -40ml3 fwd. ET from Amersham

High quality sequence stop: 256.

Location/Qualifiers

FEATURES

SOURCE

1..309

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="IMAGE:1270990"

/clone_lib="NCI_CGAP_GCB1"

/tissue_type="germinal center B cell"

/lab_host="DH10B"

/note="vector: pT7T3D-Pac (Pharmacia) with a modified

polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA

was prepared from human tonsillar cells enriched for

germinal center B cells by flow sorting (CD20+, IgD-),

provided by Dr. Louis M. Staudt (NCI), Dr. David Allman

(NCI) and Dr. Gerald Marti (CBER). cDNA synthesis was

primed with a Not I - oligo(dT) primer

[5'-TGTTACCACTCAATGGAGCGCGCCATATTTTTTTTTTTTTTTT-

3']. Double-stranded cDNA was ligated to Eco RI adaptors

(Pharmacia), digested with Not I and cloned into the Not I

and Eco RI sites of the modified pT7T3 vector. Library

went through one round of normalization, and was

constructed by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 95 a 52 c 55 g 107 t

ORIGIN

alignment_scores:

Quality: 39.00 Length: 12

Ratio: 4.333 Gaps: 0

Percent Similarity: 75.000 Percent Identity: 66.667

alignment_block:

US-08-653-294-18 x AA748739/rev ..

Align seg 1/1 to reverse of: AA748739 from: 1 to: 309

1 TyrArgLeuAlaIleArgArgIleLeuLeuArgTyr 12
||||| |||||::: ||::|||:::|||||
278 TACCGCAGAATGATACGACGCTGTACTGCTTCTCTAC 243

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run on: February 8, 2000, 01:29:40 ; Search time 122.56 Seconds
(without alignments)
2.319 Million cell updates/sec

Title: US-08-653-294-19
Perfect score: 54
Sequence: 1 YRLAIXRIALRY 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues
Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : A_Geneseq_36.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	52	96.3	12	R95429	HLA-B2702 84-79-84
2	52	96.3	12	W33798	Peptide B2702 84-7
3	52	96.3	12	W33799	Immunomodulating d
4	38	70.4	20	R92907	HLA-B2702 CTL modu
5	38	70.4	20	R95428	Immunomodulating d
6	38	70.4	20	W33778	HLA-B2702 CTL modu
7	33	61.1	20	R92908	HLA-B2702 CTL modu
8	33	61.1	20	R92908	HLA-B2702 CTL modu
9	33	61.1	20	W33791	Peptide B2702 84-7
10	33	61.1	20	W33792	Peptide B2702 84-7
11	32	59.3	11	R81449	Hepatitis GB virus
12	31	57.4	20	R95430	HLA-B2702 84-75T/7
13	29	53.7	6	W47261	Immunomodulatory p
14	29	53.7	6	W33780	Peptide #1 used in
15	29	53.7	10	R41208	Peptide fragment o
16	29	53.7	10	R83062	HLA-B2702 CTL modu
17	29	53.7	10	R83094	HLA-B2702 CTL modu
18	29	53.7	10	R95413	Alpha1-helix of HL
19	29	53.7	10	R95425	HLA-B2702.75-84(D)
20	29	53.7	10	W07512	T-cell modulating
21	29	53.7	10	W07513	T-cell modulating
22	29	53.7	10	W47265	Immunomodulatory p
23	29	53.7	10	W47269	Immunomodulatory p
24	29	53.7	10	W33784	Peptide B2702.75-8
25	29	53.7	10	W33787	Peptide B2702.75-8
26	29	53.7	15	R92912	HLA-B2702 CTL modu
27	29	53.7	15	W33795	Peptide B2702.70-8
28	29	53.7	18	R71429	Human MHC 1 alpha
29	29	53.7	25	R41205	Peptide fragment o
30	29	53.7	25	R48286	Peptide fragment o
31	29	53.7	25	R83090	HLA-B2702 CTL modu
32	29	53.7	25	R83093	HLAB38 CTL modulat
33	29	53.7	25	R95416	HLA-B2702 60-84. C
34	29	53.7	25	R95422	HLAB38.6084. Comps

Peptide B2702.60-8
Peptide Seq ID No:
Flea cysteine prot
Flea cysteine prot
Flea cysteine prot
Sequence of HLA-B5
Sequence of HLA-B5
HLA-Bw53 exon. HLA
Human neuronal nic
Recombinant cold-z
E. histolytica Pyr

ALIGNMENTS

RESULT 1

R95429 standard; peptide; 12 AA.
AC R95429;
DT 12-NOV-1996 (first entry)
DE HLA-B2702 84-79-84 palindromic.
KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN WO9513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; UI2985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Kreisky AM;
DR WPI; 95-194027/25.
PT Comps. comprising lymphoid surface membrane proteins - which may
inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 12; 29pp; English.
CC R95413, and R95431 represent palindromes and fragments of
human-leucocyte-associated antigens. This sequence represents the
HLA-B2702 84-79-84 palindromic. These sequences can be used to isolate
the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
protein associated with T-cell activation in mammalian T-cells, and is
also immunologically cross reactive with the heat shock protein Hsc70.
p74 is found in a limited number of cell types, but is particularly
expressed on B and T cells. p74 can be isolated by lysis of a suitable
cell with an amphoteric detergent, and then passed through an affinity
column containing a covalently bound HLA-B2702 palindromic peptide.
Compositions comprising the extracellular fragment of p74 combined with
HLA-B2702.60-84 (see R95416), induces calcium influx, and inhibits
cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
compounds can be screened for their effect on the cytolytic activity of
T-cells, by combining them with the extracellular portion of p74 and
determining the amount of binding between the candidate compound and p74.
Modulation of CTL activity can be inhibited in a cellular composition
containing T-cells and antigen presenting cells (APCs), by adding to the
mix the extracellular portion of p74, in an amount sufficient to compete
with p74 for the binding of the p74 ligand.
Sequence 12 AA;

Query Match 96.3%; Score 52; DB 1; Length 12;

Best Local Similarity 91.7%; Pred. No. 0.00017;

Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIXRIALRY 12

Db 1 YRLAIXRIALRY 12

RESULT 2

W33798

ID W33798 standard; peptide; 12 AA.

AC W33798;

DT 19-JUN-1998 (first entry)
 DE Peptide B2702.84-79/79-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplanted; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33799-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 12 AA;

 Query Match 96.3%; Score 52; DB 1; Length 12;
 Best Local Similarity 91.7%; Pred. No. 0.00017;
 Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

 Qy 1 YRLAIXRIALRY 12
 Db | | | | | | | | | |
 1 YRLAIRRIALRY 12

 RESULT 3
 ID W33799 standard; peptide; 12 AA.
 AC W33799;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #3.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplanted; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 17; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is

CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alpha domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 12 AA;

 Query Match 96.3%; Score 52; DB 1; Length 12;
 Best Local Similarity 91.7%; Pred. No. 0.00017;
 Matches 11; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

 Qy 1 YRLAIXRIALRY 12
 Db | | | | | | | | | |
 1 YRLAIRRIALRY 12

 RESULT 4
 ID R92907 standard; peptide; 20 AA.
 AC R92907;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.84-75/75-84).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN WO9526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI: 95-358582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host
 PS Example 15; Page 36; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

 Query Match 70.4%; Score 38; DB 1; Length 20;
 Best Local Similarity 55.0%; Pred. No. 0.16;
 Matches 11; Conservative 0; Mismatches 1; Indels 8; Gaps 1;

 Qy 1 YRLAI-----XRIALRY 12
 Db | | | | | | | | | |
 1 YRLAIRNERRRIALRY 20

RESULT 5
R95428
ID R95428 standard; peptide; 20 AA.
AC R95428;
DT 12-NOV-1996 (first entry)
DE HLA-B*2702 84-75-84 palindromic.
KW HLA: p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN W09513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 95-194027/25.
PT Comps. comprising lymphoid surface membrane proteins - which may
PT inhibit cytolytic activity and differentiation of CTLs.
PS Example: Page 12; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
CC human-leucocyte-associated antigens. These sequences can be used to isolate
CC the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
CC protein associated with T-cell activation in mammalian T-cells, and is
CC also immunologically cross reactive with the heat shock protein Hsc70.
CC p74 is found in a limited number of cell types, but is particularly
CC expressed on B and T cells. p74 can be isolated by lysis of a suitable
CC cell with an amphoteric detergent, and then passed through an affinity
CC column containing a covalently bound HLA-B*2702 palindromic peptide.
CC Compositions comprising the extracellular fragment of p74 combined with
CC HLA-B*2702.60-84 (see R95416), induces calcium influx, and inhibits
CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
CC compounds can be screened for their effect on the cytolytic activity of
CC T-cells, by combining them with the extracellular portion of p74 and
CC determining the amount of binding between the candidate compound and p74.
CC Modulation of CTL activity can be inhibited in a cellular composition
CC containing T-cells and antigen presenting cells (APCs), by adding to the
CC mix the extracellular portion of p74, in an amount sufficient to compete
CC with p74 for the binding of the p74 ligand.
CC Sequence 20 AA;

Query Match 70.4%; Score 38; DB 1; Length 20;
Best Local Similarity 55.0%; Pred. No. 0.16;
Matches 11; Conservative 0; Mismatches 1; Indels 8; Gaps 1;

QY 1 YRLAI-----XRLALRY 12
| | | | | | | | | |
DB 1 YRLAI LRLNERENLRALRY 20

RESULT 6
W33778
ID W33778 standard; peptide; 20 AA.
AC W33778;
DT 19-JUN-1998 (first entry)
DE Human immunomodulating dimer peptide #1.
KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
KW rejection.
OS Synthetic.
OS Homo sapiens.
PN W09744351-A1.
PD 27-NOV-1997; U08689.
PF 22-MAY-1997; US-653294.
PR 24-MAY-1996; US-653294.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Beulow R, Clayberger C, Krensky AM;
DR WPI: 98-086530/08.
PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
PT alpha-1 domain, used for preventing rejection of transplants or

PT treating autoimmune diseases
PS Claim 16; Page 35; 41pp; English.
CC This sequence represents a specifically claimed immunomodulating
CC dimer peptide of the invention. A peptide-type compound or variant is
CC claimed which has immunomodulating activity, including the N-terminal
CC acylated and/or C-terminal amidated or esterified forms of up to 60
CC amino acids, where the peptide-type compound comprises the formula: A-B,
CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
CC represents amino acid. The sequence in the brackets may optionally be
CC absent or truncated at any peptide type bond within the brackets. The
CC compounds comprise amino acid sequences related to a Class I HLA-B
CC alpha1 domain (positions 79-84). They can be used to inhibit cytotoxic
CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
CC vitro. They can also be used in combination with antigenic peptides or
CC proteins of interest to activate CTLs. They can also inhibit the
CC proliferation of T cells in response to anti-CD3. The peptide can be
CC used for preventing rejection of transplants or for treating autoimmune
CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
CC The products can also be used for detection and diagnosis.
CC Sequence 20 AA;

Query Match 70.4%; Score 38; DB 1; Length 20;
Best Local Similarity 55.0%; Pred. No. 0.16;
Matches 11; Conservative 0; Mismatches 1; Indels 8; Gaps 1;

QY 1 YRLAI-----XRLALRY 12
| | | | | | | | | |
DB 1 YRLAI LRLNERENLRALRY 20

RESULT 7
R92909
ID R92909 standard; peptide; 20 AA.
AC R92909;
DT 16-MAY-1996 (first entry)
DE HLA-B*2702 CTL modulating peptide (B2702.84-75/75-84(T)).
KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW immunosuppressant; graft versus host disorder; transplantation; therapy;
KW Class I MHC; HLA-B*2702.
OS Synthetic.
PN W09526979-A1.
PD 12-OCT-1995.
PF 05-APR-1995; U04349.
PR 05-APR-1994; US-222851.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM, Parham P;
DR WPI: 95-358582/46.
PT Extension of acceptance period of transplants from MHC unmatched
PT donor hosts - using Class I B75-84 MHC antigen of the recipient
PT host
PS Example 15; Page 36; 80pp; English.
CC R83061-R83085, R83090-R83096 and R92907-R92913 represent fragments of
CC class I major histocompatibility complex (MHC) antigens. This sequence
CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
CC I MHC HLA-B*2702. These sequences can be used to extend the period of
CC acceptance by a recipient of a transplant from an MHC unmatched donor.
CC The peptides are administered to a patient in conjunction with a
CC subtherapeutic amount of an immunosuppressant. This is administered to
CC the patient for a limited period of time (compared to the lifetime
CC administration for current treatments). The peptides particularly
CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
CC of the patient.
CC Sequence 20 AA;

Query Match 61.1%; Score 33; DB 1; Length 20;
Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 10; Conservative 0; Mismatches 2; Indels 8; Gaps 1;

QY 1 YRLAI-----XRLALRY 12

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Db      1 YRLAIRLNERENLRALRY 20
      |||||
      1 YRLAIRLNERENLRALRY 20

RESULT 8
R92908
ID R92908 standard: peptide; 20 AA.
AC W33791;
DT 16-MAY-1996 (first entry)
DE HLA-B*2702 CTL modulating peptide (B2702.84-75(T)/75-84).
KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
KW immunosuppressant; graft versus host disorder; transplantation; therapy;
KW Class I MHC; HLA-B*2702.
OS Synthetic.
PN WO9526979-A1.
PD 12-OCT-1995.
PF 05-APR-1995; U04349.
PR 03-APR-1994; US-222851.
PA (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM, Parham P;
DR WPI; 95-358582/46.
PT Extension of acceptance period of transplants from MHC unmatched
PT donor hosts - using Class I B75-84 MHC antigen of the recipient
PT host
PS Example 15; Page 36; 80pp; English.
CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
CC class I major histocompatibility complex (MHC) antigens. This sequence
CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
CC I MHC HLA-B*2702. These sequences can be used to extend the period of
CC acceptance by a recipient of a transplant from an MHC unmatched donor.
CC The peptides are administered to a patient in conjunction with a
CC subtherapeutic amount of an immunosuppressant. This is administered to
CC the patient for a limited period of time (compared to the lifetime
CC administration for current treatments). The peptides particularly
CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
CC of the patient.
SQ Sequence 20 AA;

Query Match 61.1%; Score 33; DB 1; Length 20;
Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 10; Conservative 0; Mismatches 2; Indels 8; Gaps 1;

QY 1 YRLAI-----XRLALRY 12
      |||||
Db      1 YRLATRLNERENLRALRY 20
      |||||

RESULT 10
W33792
ID W33792 standard; peptide; 20 AA.
AC W33791;
DT 19-JUN-1998 (first entry)
DE Peptide B2702.84-75(T)/75-84T tested for immunomodulating activity.
KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
KW rejection.
OS Synthetic.
OS Homo sapiens.
PN WO9744351-A1.
PD 27-NOV-1997.
PF 22-MAY-1997; U08689.
PR 24-MAY-1996; US-653294.
PA (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Beulow R, Clayberger C, Krensky AM;
DR WPI; 98-086530/08.
PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
PT alpha-1 domain, used for preventing rejection of transplants or
PT treating autoimmune diseases
PS Example 1; Page 19; 41pp; English.
CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
CC activity. A peptide-type compound or variant is claimed which has
CC immunomodulating activity, including the N-terminal acylated and/or
CC C-terminal amidated or esterified forms of up to 60 amino acids, where
CC the peptide-type compound comprises the formula: A-B, where A =
CC (R aa76-77L; aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
CC acid. The sequence in the brackets may optionally be absent or truncated
CC at any peptide type bond within the brackets. The compounds comprise
CC amino acid sequences related to a Class I HLA-B alpha1 domain (positions
CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
CC undesirably attacking cells in a host or in vitro. They can also be
CC used in combination with antigenic peptides or proteins of interest to
CC activate CTLs. They can also inhibit the proliferation of T cells in
CC response to anti-CD3. The peptide can be used for preventing rejection
CC of transplants or for treating autoimmune diseases, e.g. diabetes,
CC rheumatoid arthritis and lupus erythematosus. The products can also be
CC used for detection and diagnosis.
SQ Sequence 20 AA;

Query Match 61.1%; Score 33; DB 1; Length 20;
Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 10; Conservative 0; Mismatches 2; Indels 8; Gaps 1;

QY 1 YRLAI-----XRLALRY 12
      |||||
Db      1 YRLATRLNERENLRALRY 20
      |||||

RESULT 10
W33792
ID W33792 standard; peptide; 20 AA.
AC W33791;
DT 19-JUN-1998 (first entry)
DE Peptide B2702.84-75(T)/75-84T tested for immunomodulating activity.
KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
KW rejection.
OS Synthetic.
OS Homo sapiens.
PN WO9744351-A1.
PD 27-NOV-1997.
PF 22-MAY-1997; U08689.
PR 24-MAY-1996; US-653294.
PA (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Beulow R, Clayberger C, Krensky AM;
DR WPI; 98-086530/08.
PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
PT alpha-1 domain, used for preventing rejection of transplants or
PT treating autoimmune diseases
PS Example 1; Page 19; 41pp; English.
CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
CC activity. A peptide-type compound or variant is claimed which has
CC immunomodulating activity, including the N-terminal acylated and/or
CC C-terminal amidated or esterified forms of up to 60 amino acids, where
CC the peptide-type compound comprises the formula: A-B, where A =
CC (R aa76-77L; aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
CC acid. The sequence in the brackets may optionally be absent or truncated
CC at any peptide type bond within the brackets. The compounds comprise
CC amino acid sequences related to a Class I HLA-B alpha1 domain (positions
CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
CC undesirably attacking cells in a host or in vitro. They can also be
CC used in combination with antigenic peptides or proteins of interest to
CC activate CTLs. They can also inhibit the proliferation of T cells in
CC response to anti-CD3. The peptide can be used for preventing rejection
CC of transplants or for treating autoimmune diseases, e.g. diabetes,
CC rheumatoid arthritis and lupus erythematosus. The products can also be
CC used for detection and diagnosis.
SQ Sequence 20 AA;

Query Match 61.1%; Score 33; DB 1; Length 20;
Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 10; Conservative 0; Mismatches 2; Indels 8; Gaps 1;

QY 1 YRLAI-----XRLALRY 12
      |||||
Db      1 YRLATRLNERENLRALRY 20
      |||||

RESULT 10
W33791
ID W33791 standard; peptide; 20 AA.
AC W33791;
DT 19-JUN-1998 (first entry)
DE Peptide B2702.84-75(T)/75-84 tested for immunomodulating activity.
KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
KW rejection.
OS Synthetic.
OS Homo sapiens.
PN WO9744351-A1.
PD 27-NOV-1997.
PF 22-MAY-1997; U08689.
PR 24-MAY-1996; US-653294.
PA (STRD ) UNIV LELAND STANFORD JUNIOR.
PI Beulow R, Clayberger C, Krensky AM;
DR WPI; 98-086530/08.
PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
PT alpha-1 domain, used for preventing rejection of transplants or
PT treating autoimmune diseases
PS Example 1; Page 19; 41pp; English.
CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
CC activity. A peptide-type compound or variant is claimed which has
CC immunomodulating activity, including the N-terminal acylated and/or

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Best Local Similarity 50.0%; Pred. No. 1.6;
Matches 10; Conservative 0; Mismatches 2; Indels 8; Gaps 1;

QY 1 YRLAI-----XRIALRY 12
||||| |
Db 1 YRLAIRNRENLRTALRY 20

RESULT 11

R81449 ID R81449 standard; Protein: 111 AA.
AC R81449;
DE Hepatitis GB virus (HGBV) clone 50 protein prod.
KW Hepatitis GB virus; HGBV; diagnosis; treatment; vaccine;
KW reagents; non-A; non-B; non-C; non-D; non-E; clone 50;
KW tamatin; infected plasma; lambda phage; cDNA library.
OS Hepatitis GB virus
FH Key Location/Qualifiers
FT misc_difference 46 /note= "corresponding codon STOP codon"
FT misc_difference 67 /note= "corresponding codon STOP codon"
FT misc_difference 111 /note= "corresponding codon STOP codon"
FT misc_difference 111 /note= "corresponding codon STOP codon"
PN W09521922-A2.
PD 17-AUG-1995.
PF 14-FEB-1995; U02118.
PR 14-FEB-1994; US-196030.
PR 13-MAY-1994; US-242654.
PR 29-JUL-1994; US-283314.
PR 23-NOV-1994; US-344190.
PR 23-NOV-1994; US-344185.
PR 27-JAN-1995; US-344557.
PA (ABBO) ABBOTT LAB.
PI Buljk SL, Dawson GJ, Desai SM, Erker JC, Leary TP;
PI Muerhoff AS, Mushahwar IK, Pilot-Matias TJ, Schlauder GG;
PI Simons JN;
DR WPI: 95-293323/38.
DR N-PSDB: T00049.
PT Non-A, non-B, non-C, non-D, non-E Hepatitis virus reagents - useful
PT for diagnosis and therapy of hepatitis GB virus
PS Example 5; Page 22; 66pp; English.
CC Double stranded hepatitis GB virus (HGBV) DNA obtd. from HGBV
CC infected tamarin plasma, using standard procedures, was used to
CC prepare a lambda phage HGBV cDNA library. The cDNA clone T00049,
CC which encodes the proteins R81447-50 and R82064/65 (the 6 possible
CC reading frames), was rescued from the lambda phage, searched
CC against a sequence database and found to be an unique HGBV
CC sequence. Reagents which comprise the HGBV DNA, or its protein
CC prods. can be used for the diagnosis, therapy or in a vaccine to
CC prevent HGBV infection.
SQ Sequence 111 AA;

Query Match 59.3%; Score 32; DB 1; Length 111;
Best Local Similarity 41.7%; Pred. No. 13;
Matches 5; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIXRIALRY 12
| | | | | | | | | |
Db 81 YLLRMSRVAIKY 92

RESULT 12

R95430 ID R95430 standard; peptide; 20 AA.
AC R95430;
DE HLA-B2702 84-75T/75-84T palindromic.
KW HLA-B2702 84-75T/75-84T palindromic.
DE HLA; p74; alpha-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;

KW cytolysis; antigen presenting cell.
OS Synthetic.
PN W09513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 95-194027/25.
PT Compns. comprising lymphoid surface membrane proteins - which may
PT inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 12; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
CC human-leucocyte-associated antigens. This sequence represents the
CC HLA-B2702 84-75T/75-84T palindromic. These sequences can be used to
CC isolate the protein p74 from a T-cell lysate. p74 is a T-cell surface
CC membrane protein associated with T-cell activation in mammalian T-cells,
CC and is also immunologically cross reactive with the heat shock protein
CC Hsc70. p74 is found in a limited number of cell types, but is
CC particularly expressed on B and T cells. p74 can be isolated by lysis of
CC a suitable cell with an amphoteric detergent, and then passed through an
CC affinity column containing a covalently bound HLA-B2702 palindromic
CC peptide. Compositions comprising the extracellular fragment of p74
CC combined with HLA-B2702-60-84 (see R95416), induces calcium influx, and
CC inhibits cytotoxic T lymphocyte (CTL) differentiation or cytotoxicity.
CC Candidate compounds can be screened for their effect on the cytolytic
CC activity of T-cells, by combining them with the extracellular portion of
CC p74 and determining the amount of binding between the candidate compound
CC and p74. Modulation of CTL activity can be inhibited in a cellular
CC composition containing T-cells and antigen presenting cells (APCs), by
CC adding to the mix the extracellular portion of p74, in an amount
CC sufficient to compete with p74 for the binding of the p74 ligand.
SQ Sequence 20 AA;

Query Match 57.4%; Score 31; DB 1; Length 20;
Best Local Similarity 52.6%; Pred. No. 3.8;
Matches 10; Conservative 0; Mismatches 1; Indels 8; Gaps 1;

QY 1 YRLAI-----XRIALR 11
||||| |
Db 1 YRLAIRNRENLRIALR 19

RESULT 13
W47261 ID W47261 standard; peptide; 6 AA.
AC W47261;
DE 22-MAY-1998 (first entry)
DE Immunomodulatory peptide.
KW Immunomodulator; Class I HLA-B alpha-1 domain; inhibition;
KW transplant rejection; treatment; autoimmune disease.
OS Homo sapiens.
OS Synthetic.
FH Key Location/Qualifiers

FT Misc_difference 1. .6 /note= "at least one of the amino acids is the
FT D-isomer

PN W09744052-A1.
PD 27-NOV-1997.
PF 23-APR-1997; U06705.
PR 22-MAY-1996; US-651650.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 98-018220/02.
PT Novel immunomodulatory peptide-type compound - useful for inhibiting
PT transplant rejection

PS Claim 10; Page 36; 41pp; English.
CC The present sequence is an immunomodulatory peptide, which
CC comprises a Class I HLA-B alpha-1 domain sequence. It can be used
CC in a pharmaceutical composition together with a subtherapeutic dose
CC of an immunosuppressant, to extend the period of acceptance of a
CC transplant from a major histocompatibility complex (MHC) unmatched

CC donor, i.e. to inhibit transplant rejection. It can also be used in
 CC the treatment of autoimmune diseases.
 CC Peptides using the D-form amino acids are more effective
 CC immunomodulators than their diastereomers or enantiomers.
 SQ Sequence 5 AA;

Query Match 53.7%; Score 29; DB 1; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.5e+05;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 RIALRY 12
 |||||
 DB 1 RIALRY 6

RESULT 14

W33780
 ID W33780 standard; peptide: 6 AA.
 AC W33780;
 DT 19-JUN-1998 (first entry)
 DE Peptide #1 used in immunomodulating dimer peptide.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN WO9744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 WPI: 98-086530/08.

PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 15; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed peptide which forms part
 CC of the immunomodulating dimer peptides of the invention. A peptide-type
 CC compound or variant is claimed which has immunomodulating activity,
 CC including the N-terminal acylated and/or C-terminal amidated or
 CC esterified forms of up to 60 amino acids, where the peptide-type compound
 CC comprises the formula: A-B, where A, B = (R aa76-77L) (aa79-84) or
 CC (aa84-79) (Laa77-76R); aa76 = E or V; aa77 = D, S or N; aa79 = R or G;
 CC aa80 = I or N; aa81, aa84 = a hydrophobic or small amino acid; aa82 = R
 CC or L; aa83 = G or R; and aa represents amino acid. The sequence in the
 CC brackets may optionally be absent or truncated at any peptide type bond
 CC within the brackets. The compounds comprise amino acid sequences related
 CC to a Class I HLA-B alpha1 domain (positions 79-84). They can be used to
 CC inhibit cytotoxic T-lymphocytes (CTL) from undesirably attacking cells in
 CC a host or in vitro. They can also be used in combination with antigenic
 CC peptides or proteins of interest to activate CTLs. They can also inhibit
 CC the proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 6 AA;

Query Match 53.7%; Score 29; DB 1; Length 6;
 Best Local Similarity 100.0%; Pred. No. 1.5e+05;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 RIALRY 12
 |||||
 DB 1 RIALRY 6

RESULT 15

R41208
 ID R41208 standard; peptide: 10 AA.
 AC R41208;

DT 15-MAR-1994 (first entry)
 DE Peptide fragment of Class I HLA peptide.
 KW Human leukocyte antigen; HLA; peptide; transplantation; neoplasia;
 KW parasitic disease; cytotoxic T lymphocyte; modulation.
 OS Synthetic.
 PN WO9317699-A.
 PD 16-SEP-1993.
 PF 25-FEB-1993; U01758.
 PR 02-MAR-1992; US-844716.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger CA, Krensky AM;
 DR WPI: 93-303134/38.
 DT New peptide(s) based on Class I HLA antigen domains - used for
 PT modulating cytotoxic T-lymphocyte activity towards targets
 PS Claim 11; Page 54; 61pp; English.
 CC The peptide is used to modulate cytotoxic T-lymphocyte (CTL)
 CC activity, either by inhibition or stimulation. It can be used
 CC for inhibiting CTL toxicity in transplantations, for inducing CTL
 CC activity in parasitic diseases and neoplasia and in studies on viral
 CC infection. The peptide can also be used for identifying CTLs which
 CC bind to it and removing subsets of CTLs from a T-cell composition.
 CC This peptide sequence is more commonly found within larger peptide
 CC compounds of not more than 30 amino acids in length.
 SQ Sequence 10 AA;

Query Match 53.7%; Score 29; DB 1; Length 10;
 Best Local Similarity 100.0%; Pred. No. 4.8;
 Matches 6; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 7 RIALRY 12
 |||||
 DB 5 RIALRY 10

Search completed: February 8, 2000, 01:29:40
 Job time: 1752 sec

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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:29 ; Search time 117.7 seconds
(without alignments)
4.809 Million cell updates/sec

Title: US-08-653-294-19
Perfect score: 54
Sequence: 1 YRLAIXRLRY 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : PIR62:*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	34	63.0	141	2 H71504	ribosomal protein
2	34	63.0	309	2 G70882	probable oxidoreductase
3	33	61.1	142	2 C72053	L17 ribosomal prot
4	32	59.3	253	2 E75055	hypothetical prote
5	32	59.3	264	2 E71567	hypothetical prote
6	32	59.3	552	2 A84085	probable ATP-bindi
7	32	59.3	610	2 S77337	ABC-type transport
8	32	59.3	2242	2 A57541	pyrimidine synthe
9	31	57.4	190	2 F70410	hypothetical prote
10	31	57.4	349	2 G70542	probable bioB prot
11	31	57.4	356	2 S37356	spas protein - sal
12	31	57.4	469	2 C69460	conserved hypothet
13	31	57.4	578	2 B64012	hypothetical prote
14	31	57.4	1166	2 A49201	adenylate cyclase
15	31	57.4	1180	2 A47202	adenylate cyclase
16	31	57.4	1791	2 T02909	hypothetical prote
17	30	55.6	88	2 C70200	ribosomal protein
18	30	55.6	153	2 S08501	3-dehydroquinat d
19	30	55.6	211	2 H64961	probable membrane
20	30	55.6	283	2 JC6531	avermectin B 5-O-m
21	30	55.6	286	1 B69290	riannosyl transfer
22	30	55.6	313	2 JC5342	Na+/H+ antiporter
23	30	55.6	331	2 S09800	hypothetical prote
24	30	55.6	348	2 S29990	histocompatibility
25	30	55.6	376	2 S75438	hypothetical prote
26	30	55.6	377	2 A72350	pleiotropic regula
27	30	55.6	394	2 S20905	hypothetical prote
28	30	55.6	432	2 F70575	probable Pura - My
29	30	55.6	497	2 D65169	yisc protein - Esc
30	30	55.6	528	2 B64760	propionate catabol

31 30 55.6 564 2 I64134
32 30 55.6 690 2 S54211
33 30 55.6 754 2 S52564
34 30 55.6 770 1 W2BE30
35 30 55.6 880 2 F71652
36 30 55.6 1084 2 G71329
37 30 55.6 2347 1 TVHURS
38 30 55.6 3712 2 S18253
39 29 53.7 124 1 QOVZEL
40 29 53.7 133 2 H42513
41 29 53.7 133 2 S33096
42 29 53.7 138 1 A89219
43 29 53.7 144 2 H72464
44 29 53.7 156 2 T12893
45 29 53.7 170 2 E69149

D-lactate dehydrog
ATM1 protein precu
hypothetical prote
gene 30 protein -
pyruvate,phosphate
hypothetical prote
kinase-related pro
laminin alpha-1 ch
F11 protein - vacc
J5L protein - vacc
J5L protein - vari
conserved hypothet
hypothetical prote
hypothetical prote
hypothetical prote

ALIGNMENTS

RESULT 1
H71504
ribosomal protein L17 - Chlamydia trachomatis
C:Species: Chlamydia trachomatis
C:Date: 13-Sep-1998 #sequence_revision 13-Sep-1998 #text_change 13-Aug-1999
C:Accession: H71504; I40747
R:Stephens, R.S.; Kalman, S.; Lammel, C.J.; Fan, J.; Marathe, R.; Aravind, L.; Mitche
Science 282, 754-759, 1998
A:Title: Genome sequence of an obligate intracellular pathogen of humans: Chlamydia t
A:Reference number: A71570; MUID:99000809
A:Accession: H71504
A:Molecule type: DNA
A:Residues: 1-141 <ARN>
A:Cross-references: GB:AE001323; GB:AE001273; NID:g3328931; PIDN:AAC68107.1; PID:g332
A:Experimental source: serotype D, strain UW-3/CX
R:Gu, L.; Wenman, W.M.; Remacha, M.; Meuser, R.; Coffin, J.; Kaul, R.
J. Bacteriol. 177, 2594-2601, 1995
A:Title: Chlamydia trachomatis RNA polymerase alpha subunit: sequence and structural
A:Reference number: I40743; MUID:95247702
A:Accession: I40747
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 20-115, 'R', 117-141 <GUL>
A:Cross-references: GB:L33834; NID:g620026; PIDN:AAA74990.1; PID:g620030
C:Genetics:
A:Gene: r117
C:Superfamily: Escherichia coli ribosomal protein L17
C:Keywords: protein biosynthesis; ribosome

Query Match 63.0%; Score 34; DB 2; Length 141;
Best Local Similarity 54.5%; Pred. No. 3.7;
Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIIXRLRY 12
|||:|:|:
DB 64 RLAVGLMVRV 74

RESULT 2
G70882
probable oxidoreductase - Mycobacterium tuberculosis (strain H37RV)
C:Species: Mycobacterium tuberculosis
C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 29-Sep-1999
C:Accession: G70882
R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon
; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd,
Nature 393, 537-544, 1998
A:Authors: Krogh, A.; McLean, J.; Moule, S.; Murphy, L.; Oliver, S.; Osborne, J.; Qua
; Taylor, K.; Whitehead, S.; Barrell, B.G.
A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno
A:Reference number: A70500; MUID:98295987
A:Accession: G70882

A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-309 <COL>
A:Cross-references: GB:AL008967; GB:AL123456; NID:g3261491; PIDN:CAAL5591.1; PID:e129972
A:Experimental source: strain H37Rv
C:Genetics:
A:Gene: RV2776c
C:Superfamily: phthalate dioxygenase reductase; cytochrome-b5 reductase homology; ferredoxin
F:9-211/Domain: cytochrome-b5 reductase homology <CBR>
F:240-297/Domain: ferredoxin [2Fe-2S] homology <FER>

Query Match 63.0%; Score 34; DB 2; Length 309;
Best Local Similarity 77.8%; Pred. No. 8;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIXRIA 9
||| |||
DB 65 YRLAIRRIA 73

RESULT 3
C72053
L17 ribosomal protein - Chlamydia pneumoniae (strain CWL029)
C:Species: Chlamydia pneumoniae
C:Date: 23-Apr-1999 #sequence_revision 23-Apr-1999 #text_change 13-Aug-1999
C:Accession: C72053
R:Kallman, S.; Mitchell, W.; Marathe, R.; Lammel, C.; Fan, J.; Olinger, L.; Grimwood, J.; Nature Genet. 21, 385-389, 1999
A:title: Comparative genomes of Chlamydia pneumoniae and C. trachomatis.
A:Reference number: A72000; MUID:99206606
A:Accession: C72053
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-142 <ARN>
A:Cross-references: GB:AE001647; GB:AE001363; NID:g4376920; PIDN:AAD18764.1; PID:g437692
A:Experimental source: strain CWL029
C:Genetics:
A:Gene: r117
C:Superfamily: Escherichia coli ribosomal protein L17

Query Match 61.1%; Score 33; DB 2; Length 142;
Best Local Similarity 54.5%; Pred. No. 6.1;
Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIIXRIALRY 12
||| |||
DB 64 RLAIIRLMVRY 74

RESULT 4
E75055
Hypothetical protein PAB1428 - Pyrococcus abyssi (strain Orsay)
C:Species: Pyrococcus abyssi
C:Date: 20-Aug-1999 #sequence_revision 20-Aug-1999 #text_change 20-Aug-1999
C:Accession: E75055
R:anonymous, Genoscope
submitted to the EMBL Data Library, July 1999
A:Description: Pyrococcus abyssi genome sequence: insights into archaeal chromosome structure
A:Reference number: A75001
A:Accession: E75055
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-253 <KAW>
A:Cross-references: GB:AJ248287; GB:AL096836; NID:g5458657; PIDN:CAB50338.1; PID:e151623
A:Experimental source: strain Orsay
C:Genetics:
A:Gene: PAB1428

Query Match 59.3%; Score 32; DB 2; Length 253;
Best Local Similarity 66.7%; Pred. No. 18;

A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-309 <COL>
A:Cross-references: GB:AL008967; GB:AL123456; NID:g3261491; PIDN:CAAL5591.1; PID:e129972
A:Experimental source: strain H37Rv
C:Genetics:
A:Gene: RV2776c
C:Superfamily: phthalate dioxygenase reductase; cytochrome-b5 reductase homology; ferredoxin
F:9-211/Domain: cytochrome-b5 reductase homology <CBR>
F:240-297/Domain: ferredoxin [2Fe-2S] homology <FER>

Query Match 59.3%; Score 32; DB 2; Length 264;
Best Local Similarity 70.0%; Pred. No. 18;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIIXRIALRY 11
||| |||
DB 83 RLALSRIALRY 92

RESULT 6
A64085
Probable ATP-binding transport protein HI0664 - Haemophilus influenzae (strain Rd KW2)
C:Species: Haemophilus influenzae
C:Date: 18-Aug-1995 #sequence_revision 18-Aug-1995 #text_change 20-Sep-1999
C:Accession: A64085
R:Fleischmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage
; Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman
, D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghagen, N.S.M.
Science 269, 496-512, 1995
A:Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter
A:title: Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.
A:Reference number: A64000; MUID:95350630
A:Accession: A64085
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-552 <TIGR>
A:Cross-references: GB:U32749; GB:L42023; NID:g1573658; PIDN:AC22321.1; PID:g1573663
C:Superfamily: unassigned ATP-binding cassette proteins; ATP-binding cassette homology
C:Keywords: ATP; P-loop
F:355-549/Domain: ATP-binding cassette homology <ABC>
F:372-379/Region: nucleotide-binding motif A (P-loop)

Query Match 59.3%; Score 32; DB 2; Length 552;
Best Local Similarity 70.0%; Pred. No. 38;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 3 LAIXRIALRY 12
||| |||
DB 70 LAVARGALRY 79

RESULT 7
S77337
ABC-type transport protein sll1725 - Synechocystis sp. (strain PCC 6803)

N:Alternate names: ABC transporter; protein sl11725
C:Species: *Synechocystis* sp.

A:Variety: PCC 6803
C:Date: 25-Apr-1997 #sequence_revision 25-Apr-1997 #text_change 20-Sep-1999

C:Accession: S77337
R:Kaneko, T.; Sato, S.; Kotani, H.; Tanaka, A.; Asamizu, Y.; Miyajima, N.; O. K.; Okumura, S.; Shimpo, S.; Takeuchi, C.; Wada, T.; Watanabe, A.; Yamada, M.; Yasuda DNA Res. 3, 109-136, 1996

A:Title: Sequence analysis of the genome of the unicellular cyanobacterium *Synechocystis* sp.

A:Reference number: S74322; MUID:97061201

A:Accession: S77337

A>Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-610 <KAN>

A:Cross-references: EMBL:D90906; GB:AB001339; NID:g1652492; PIDN:BAAL7440.1; PID:d101817

A:Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1996

C:Superfamily: unassigned ATP-binding cassette proteins; ATP-binding cassette homology

C:Keywords: ATP; P-loop; transport protein

F:380-574/Domain: ATP-binding cassette homology <ABC>

F:397-404/Region: nucleotide-binding motif A (P-loop)

Query Match 59.3%; Score 32; DB 2; Length 610;

Best Local Similarity 70.0%; Pred. No. 42;

Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Qy 2 RLAIAXRIALR 11

||||| I I I

Db 508 RLAIARAAMR 517

RESULT 8

A57541

pyrimidine synthesis multifunctional protein CAD - spiny dogfish

N:Contents: aspartate carbamoyltransferase (EC 2.1.3.2); carbamoyl-phosphate synthase (9

C:Species: *Squalus acanthias* (spiny dogfish)

C:Date: 08-Feb-1996 #sequence_revision 08-Feb-1996 #text_change 22-Jun-1999

C:Accession: A57541

R:Hong, J.; Salo, W.L.; Anderson, P.M.

J. Biol. Chem. 270, 14130-14139, 1995

A:Title: Nucleotide sequence and tissue-specific expression of the multifunctional prote

A:Reference number: A57541; MUID:95294021

A:Accession: A57541

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-2242 <HON>

A:Cross-references: GB:U18668; NID:g951095; PIDN:AAA74569.1; PID:g951096

C:Superfamily: rudimentary enzyme; aspartate/ornithine carbamoyltransferase homology; Ba

carbamoyl-phosphate synthase (glutamine-hydrolyzing) large chain homology; carbamoyl-phos

C:Keywords: hydrolase; ligase; transferase

F:4-1449/Domain: carbamoyl-phosphate synthase (ammonia) homology <CPA>

F:4-354/Domain: trpG homology <TRG>

F:178-354/Domain: carbamoyl-phosphate synthase (glutamine-hydrolyzing) small chain homolog

F:398-1446/Domain: carbamoyl-phosphate synthase (glutamine-hydrolyzing) large chain hom

F:998-849/Domain: biotin carboxylase homology <BC1>

F:939-1385/Domain: biotin carboxylase homology <BC2>

F:1464-1808/Domain: *Bacillus dihydroxyotase* homology <DHO>

F:1941-2239/Domain: aspartate/ornithine carbamoyltransferase homology <ACT>

F:252/Active site: Cys #status predicted

Query Match

Best Local Similarity 59.3%; Score 32; DB 2; Length 2242;

Matches 6; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 YRLAIXRIALRY 12

| | | | |

Db 2109 YLLTLRVNRLY 2120

RESULT 9

F70410

hypothetical protein aq_1277 - Aquifex aeolicus

C:Species: *Aquifex aeolicus*

C:Date: 08-May-1998 #sequence_revision 08-May-1998 #text_change 08-May-1998

C:Accession: F70410

R:Deckert, G.; Warren, P.V.; Gaasterland, T.; Young, W.G.; Lenox, A.L.; Graham, D.E.;

Nature 392, 353-358, 1998

A:Title: The complete genome of the hyperthermophilic bacterium *Aquifex aeolicus*.

A:Reference number: A70300; MUID:98196666

A:Accession: F70410

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-190 <AQF>

A:Cross-references: GB:AE000732; NID:g2983704; PID:g2983715; GB:AE000657

A:Experimental source: strain VF5

C:Genetics:

A:Gene: aq_1277

Query Match

Best Local Similarity 57.4%; Score 31; DB 2; Length 190;

Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLAIXRIALR 11

: ||| : |||

Db 43 HNLAIQKVALR 53

RESULT 10

G70542

probable bioB protein - Mycobacterium tuberculosis (strain H37RV)

C:Species: *Mycobacterium tuberculosis*

C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 16-Jul-1999

C:Accession: G70542

R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon

Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holroyd,

Rajandram, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.

Nature 393, 537-544, 1998

A:Authors: Squares, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrell, B.G.

A:Title: Deciphering the biology of *Mycobacterium tuberculosis* from the complete geno

A:Reference number: A70500; MUID:98295987

A:Accession: G70542

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-349 <COL>

A:Cross-references: GB:295586; GB:AL123456; NID:g3261785; PIDN:CAB09080.1; PID:e31716

A:Experimental source: strain H37RV

C:Genetics:

A:Gene: bioB

C:Superfamily: biotin synthetase

Query Match

Best Local Similarity 57.4%; Score 31; DB 2; Length 349;

Matches 6; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

Qy 1 YRLAIXRIALRY 12

: ||| : |||

Db 280 FRALPRTMLRF 291

RESULT 11

S37356

spas protein - *Salmonella typhimurium*

C:Species: *Salmonella typhimurium*

C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 20-Sep-1999

C:Accession: S37356

R:Groisman, E.A.; Ochman, H.

EMBO J. 12, 3779-3787, 1993

A:Title: Cognate gene clusters govern invasion of host epithelial cells by *Salmonella*

A:Reference number: S37304; MUID:94008985

A:Accession: S37356

A>Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-356 <GRO>

A:Cross-references: EMBL:X73525; NID:g404286; PID:g404293

A>Note: the nucleotide sequence was submitted to the EMBL Data Library, June 1993

C:Genetics:

A:Gene: spaS

C:Superfamily: flagellar biosynthetic protein flhB; flhB carboxyl-terminal homology

C:Keywords: transmembrane protein

Query Match 57.4%; Score 31; DB 2; Length 356;

Best Local Similarity 50.0%; Pred. No. 40;

Matches 6; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIXRIALRY 12

Db 70 YSLAVFGIGLYK 81

RESULT 12

C69460

conserved hypothetical protein Arl684 - Archaeoglobus fulgidus

C:Species: Archaeoglobus fulgidus

C:Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 29-Sep-1999

C:Accession: C69460

R:Klenk, H.P.; Clayton, R.A.; Tomb, J.F.; White, O.; Nelson, K.E.; Ketchum, K.A.; Dodson

; Fleischmann, R.D.; Quackenbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E.F.

; Glodek, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L.

Nature 390, 364-370, 1997

A:Authors: Uterback, T.; Cotton, M.D.; Spriggs, T.; Artiach, P.; Kaine, B.P.; Sykes, S.

Smith, H.O.; Woese, C.R.; Venter, J.C.

A:Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing archae

A:Reference number: A69250; MUID:98049343

A:Accession: C69460

A>Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-469 <KLE>

A:Cross-references: GB:AE000987; GB:AE000782; NID:g2689310; PIDN:AAB89565.1; PID:g264887

C:Superfamily: hypothetical protein MJ0966

Query Match 57.4%; Score 31; DB 2; Length 469;

Best Local Similarity 58.3%; Pred. No. 53;

Matches 7; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIXRIALRY 12

Db 12 YRGMKRIALVY 23

RESULT 13

B64012

hypothetical protein HI0698 - Haemophilus influenzae (strain Rd KW20)

C:Species: Haemophilus influenzae

C:Date: 18-Aug-1995 #sequence_revision 18-Aug-1995 #text_change 10-Oct-1997

C:Accession: B64012

R:Fleischmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage, A.

; Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman, J.

; D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghagen, N.S.M.

Science 269, 496-512, 1995

A:Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter,

A:Title: Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.

A:Reference number: A64000; MUID:95350630

A:Accession: B64012

A>Status: nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-578 <TIGR>

A:Cross-references: GB:032752; GB:I42023; NID:gl573692; PID:gl573700; TIGR:HI0698

Query Match 57.4%; Score 31; DB 2; Length 578;

Best Local Similarity 60.0%; Pred. No. 65;

Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLAIXRIAL 10

Db 147 YKTAISRLAL 156

RESULT 14

A49201

adenylate cyclase (EC 4.6.1.1) type V, calcium-inhibitable - mouse

C:Species: Mus musculus (house mouse)

C:Date: 19-Dec-1993 #sequence_revision 18-Nov-1994 #text_change 16-Jul-1999

C:Accession: A49201; A46187

R:Prenont, R.T.; Jacobowitz, O.; Iyengar, R.

Endocrinology 131, 2774-2784, 1992

A:Title: Lowered responsiveness of the catalyst of adenylyl cyclase to stimulation by

A:Reference number: A49201; MUID:93076707

A:Accession: A49201

A:Molecule type: mRNA

A:Residues: 7-1161 <PRE>

A:Cross-references: GB:M96653; NID:gl91726; PIDN:AAA37182.1; PID:gl91727

A:Experimental source: S49 lymphoma cells

A>Note: sequence extracted from NCBI backbone (NCBIN:119384, NCBIP:119386)

R:Yoshimura, M.; Cooper, D.M.

Proc. Natl. Acad. Sci. U.S.A. 89, 6716-6720, 1992

A:Title: Cloning and expression of a Ca(2+)-inhibitable adenylyl cyclase from NCB-20

A:Reference number: A46187; MUID:92357702

A:Accession: A46187

A>Status: not compared with conceptual translation

A:Molecule type: mRNA

A:Residues: 1-46, 'K', 48-75, 'G', 77-507, 'GR', 511-737, 'V', 739-881, 'L', 883-990, 'V', 992-11

A:Cross-references: GB:M93422; NID:gl91690; PIDN:AAA37174.1; PID:gl91691

A:Experimental source: NCB-20 cells

A>Note: sequence extracted from NCBI backbone (NCBI:110233); the authors acknowledge

C:Superfamily: human adenylyl cyclase; guanylate cyclase catalytic domain homology

C:Keywords: phosphorus-oxygen lyase; transmembrane protein

F:319-554/Domain: guanylate cyclase catalytic domain homology <GCC>

F:925-1164/Domain: guanylate cyclase catalytic domain homology <GCC>

Query Match 57.4%; Score 31; DB 2; Length 1166;

Best Local Similarity 55.6%; Pred. No. 1.3e+02;

Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 4 AIXRIALRY 12

Db 886 AVGRVALKY 894

RESULT 15

A47202

adenylate cyclase (EC 4.6.1.1) type 6 - rat

C:Species: Rattus norvegicus (Norway rat)

C:Date: 21-Jan-1994 #sequence_revision 18-Nov-1994 #text_change 16-Jul-1999

C:Accession: A47202; A45145

R:Prenont, R.T.; Chen, J.; Ma, H.W.; Ponnappalli, M.; Iyengar, R.

Proc. Natl. Acad. Sci. U.S.A. 89, 9809-9813, 1992

A:Title: Two members of a widely expressed subfamily of hormone-stimulated adenylyl c

A:Reference number: A47202; MUID:93028552

A:Accession: A47202

A>Status: preliminary; not compared with conceptual translation

A:Molecule type: mRNA

A:Residues: 1-1180 <PRE>

A:Experimental source: liver, kidney

A>Note: sequence extracted from NCBI backbone (NCBI:115851)

R:Krupinski, J.; Lehman, T.C.; Frankfield, C.D.; Zwaagstra, J.C.; Watson, P.A.

J. Biol. Chem. 267, 24858-24862, 1992

A:Title: Molecular diversity in the adenylyl cyclase family. Evidence for eight forms

A:Reference number: A45145; MUID:93077589

A:Accession: A45145

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 15-93, 'K', 95-463, 'V', 465-551, 'G', 553-803, 'I', 805-1180 <KRU>

A:Cross-references: GB:I01115; NID:g202712; PIDN:AAA40676.1; PID:g202713

C:Superfamily: human adenylate cyclase; guanylate cyclase catalytic domain homology
C:Keywords: phosphorus-oxygen lyase
F:333-568/Domain: guanylate cyclase catalytic domain homology <GCC>
F:939-1178/Domain: guanylate cyclase catalytic domain homology <GCC2>

Query Match 57.4%; Score 31; DB 2; Length 1180;
Best Local Similarity 55.6%; Pred. No. 1.3e+02;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 4 AIXRIALRY 12
I: |.:|.:|
Db 900 AVGRVALKY 908

Search completed: February 7, 2000, 11:54:30
Job time: 24340 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 00:59:57 ; Search time 63.71 Seconds
(without alignments)
5.625 Million cell updates/sec

Title: US-08-653-294-19
Perfect score: 54
Sequence: 1 YRLAIXRIALRY 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 82229 seqs, 29864866 residues

Total number of hits satisfying chosen parameters: 82229

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : SwissProt_38.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	34	63.0	122	1 RL17_CHLTR	P4760 chlamydia t
2	33	61.1	375	1 HRMA_PSESY	Q08370 pseudomonas
3	32	59.3	552	1 Y664_HAEIN	Q57538 haemophilus
4	32	59.3	2242	1 PRL1_SQUAC	Q91437 squalus aca
5	31	57.4	349	1 B10B_MYCTU	O06601 mycobacteri
6	31	57.4	356	1 SPAS_SALTY	P40702 salmonella
7	31	57.4	456	1 YD45_SCHPO	Q10299 schizosacch
8	31	57.4	541	1 PRPR_SALTY	P74839 salmonella
9	31	57.4	578	1 YTFM_HAEIN	P44038 haemophilus
10	31	57.4	582	1 MNT_HUMAN	Q99583 homo sapien
11	31	57.4	591	1 MNT_MOUSE	O08789 mus musculu
12	31	57.4	833	1 YPD2_CAEEL	Q11182 caenorhabdi
13	31	57.4	1165	1 CYA6_MOUSE	Q01341 mus musculu
14	31	57.4	1166	1 CYA6_RAT	Q03343 rattus norv
15	31	57.4	2569	1 LNA3_MOUSE	Q61789 mus musculu
16	30	55.6	88	1 RS15_BORBU	Q51744 borrelia bu
17	30	55.6	153	1 3DHQ_EMENI	P05147 emeritella
18	30	55.6	331	1 UL38_HCMVA	P16779 human cytom
19	30	55.6	348	1 HFAF_MACMU	P33617 macaca mula
20	30	55.6	432	1 PURA_MYCTU	O08381 mycobacteri
21	30	55.6	497	1 YIGC_ECOLI	P26615 escherichia
22	30	55.6	528	1 PRPR_ECOLI	P77743 escherichia
23	30	55.6	564	1 LDHD_HAEIN	P45295 haemophilus
24	30	55.6	690	1 ATM1_YEAST	P40416 saccharomyc
25	30	55.6	770	1 PRTP_VZVD	P09284 varicella-z
26	30	55.6	2347	1 KROS_HUMAN	Q08922 homo sapien
27	30	55.6	3712	1 LNA_DROME	Q00174 drosophila
28	29	53.7	124	1 VJ05_VACCV	P07618 vaccinia vi
29	29	53.7	133	1 VJ05_VACCC	P21083 vaccinia vi
30	29	53.7	133	1 VJ05_VARY	P33055 variola vir
31	29	53.7	185	1 PTH_RICPR	Q92cv4 rickettsia
32	29	53.7	203	1 NH10_YEAST	Q03435 saccharomyc
33	29	53.7	281	1 STRF_STRGR	P09397 streptomyce
34	29	53.7	337	1 CMST_HUMAN	P78382 homo sapien

ALIGNMENTS

RESULT 1
RL17_CHLTR

AC P4760; STANDARD; PRT; 122 AA.
DT 01-FEB-1996 (Rel. 33, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE 50S RIBOSOMAL PROTEIN L17.
GN RPLQ.
OS Chlamydia trachomatis.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=434/BU / SEROVAR L2;
RX MEDLINE; 95247702.
RA GU L.J., WENMAN W.M., REMACHA M., MEUSER R.U., COFFIN J.M., KAUL R.;
RT "Chlamydia trachomatis RNA polymerase alpha subunit: sequence and structural analysis."
RL J. Bacteriol. 177:2594-2601(1995).
CC -!- SIMILARITY: BELONGS TO THE L17P FAMILY OF RIBOSOMAL PROTEINS.

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CC -----
DR EMBL; L33834; AAA74990.1; .
DR PROSITE; PS01167; RIBOSOMAL_L17; 1.
DR PFAM; PF01196; Ribosomal_L17; 1.
KW Ribosomal protein.
SQ SEQUENCE 122 AA; 13969 MW; B8C43F7D CRC32;
Query Match 63.0%; Score 34; DB 1; Length 122;
Best Local Similarity 54.5%; Pred. No. 1.3; Indels 0; Gaps 0;
Matches 6; Conservative 3; Mismatches 2;

Qy 2 RLAIIXRIALRY 12
|||:|:|
Db 45 RLAVGLMVRY 55

RESULT 2
HRMA_PSESY

ID HRMA_PSESY STANDARD; PRT; 375 AA.
AC Q08370;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE HRMA PROTEIN.
GN HRMA.
OS Pseudomonas syringae (pv. syringae).
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonas group; Pseudomonas.

CC -1- MISCELLANEOUS: GATASE (GLUTAMINE AMIDOTRANSFERASE) AND CPSASE
 CC (CARBAMOYL PHOSPHATE SYNTHASE) FORM TOGETHER THE
 CC GLUTAMINE-DEPENDENT CPSASE (GD-CPSASE) (EC 6.3.5.5).
 CC -1- SIMILARITY: THE CPSASE DOMAIN IS SIMILAR TO OTHER CPASES; THE
 CC DHOASE TO OTHER DHOASES.
 CC -1- SIMILARITY: THE GATASE DOMAIN BELONGS TO TYPE-1 GLUTAMINE
 CC AMIDOTRANSFERASES.
 CC -----
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 CC -----
 CC EMBL: U18668; AAA74569.1; --
 CC HSSP: P00479; 3ATL.
 CC PROSITE: PS00097; CARBAMOYLTRANSFERASE; 1.
 CC PROSITE: PS00442; GATASE_TYPE_1; 1.
 CC PROSITE: PS00482; DIHYDROOROTASE_1; 1.
 CC PROSITE: PS00483; DIHYDROOROTASE_2; 1.
 CC PROSITE: PS00866; CPSASE_1; 2.
 CC PROSITE: PS00867; CPSASE_2; 2.
 CC PFAM: PF00117; GATase; 1.
 CC PFAM: PF00185; OTCace; 1.
 CC PFAM: PF00289; CPSase_L_chain; 2.
 CC PFAM: PF00744; Dihydroorotase; 1.
 CC PFAM: PF00988; CPSase_sm_chain; 1.
 CC Pyrimidine biosynthesis; Ligase; Transferase; Hydrolase; Zinc;
 CC Allosteric enzyme; Multifunctional enzyme; Phosphorylation.
 FT DOMAIN 1 365 GATASE (GLUTAMINE AMIDOTRANSFERASE).
 FT DOMAIN 366 397 LINKER.
 FT DOMAIN 398 1462 CPSASE (CARBAMOYL-PHOSPHATE SYNTHASE).
 FT DOMAIN 398 937 CPSASE A.
 FT DOMAIN 938 1462 CPSASE B.
 FT DOMAIN 1463 1796 DHOASE (DIHYDROOROTASE).
 FT DOMAIN 1797 1934 LINKER.
 FT DOMAIN 1935 2242 ATCase (ASPARTATE TRANSAMYLASE).
 FT ACT_SITE 252 252 GATASE (BY SIMILARITY).
 FT ACT_SITE 336 336 GATASE (BY SIMILARITY).
 FT ACT_SITE 338 338 GATASE (BY SIMILARITY).
 FT METAL 1478 1478 ZINC (POTENTIAL).
 FT METAL 1480 1480 ZINC (POTENTIAL).
 SQ SEQUENCE 2242 AA; 249391 MW; 9F38F825 CRC32;

 Query Match 59.3%; Score 32; DB 1; Length 2242;
 Best Local Similarity 50.0%; Pred. No. 84;
 Matches 6; Conservative 2; Mismatches 4; Indels 0; Gaps 0;
 QY 1 YRLAIXRIALRY 12
 | | | | |
 Db 2109 YLLTLRVNRLY 2120

 RESULT 5
 BIOB_MYCTU STANDARD; PRT; 349 AA.
 ID BIOB_MYCTU
 AC O06601;
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE BIOTIN SYNTHASE (EC 2.8.1.6) (BIOTIN SYNTHETASE).
 GN BIOB OR RV1589 OR MTCY336.15C.
 OS Mycobacterium tuberculosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=H37RV;
 RX MEDLINE: 98295987.
 RA COLE S.T., BROSCHE R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,

RA GORDON S.V., EIGLMEIER K., GAS S., BARRY C.E. III, TEKAIA F.,
 RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
 RA DAVIES R., DEVLIN K., FELTWEILL T., GENTILES S., HAMLIN N., HOLROYD S.,
 RA HORNBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,
 RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAN M.A., ROGERS J.,
 RA RUTTER S., SEGER K., SKELTON S., SQUARES S., SOARES R., SULSTON J.E.,
 RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
 RT "Deciphering the biology of Mycobacterium tuberculosis from the
 RL complete genome sequence."
 RL Nature 393:537-544(1998).
 RN SEQUENCE FROM N.A.
 RP STRAIN=PASTEUR;
 RC YU S., JACOBS W.R. JR.;
 RL Submitted (JAN-1998) to the EMBL/GenBank/DBJ databases.
 CC -1- CATALYTIC ACTIVITY: DETHIOBIOTIN + (S) - BIOTIN
 CC -1- PATHWAY: LAST STEP IN BIOTIN BIOSYNTHESIS PATHWAY.
 CC -1- SIMILARITY: BELONGS TO THE BIOTIN AND LIPOIC ACID SYNTHETASES
 CC FAMILY.
 CC -----
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 CC -----
 CC EMBL: Z95586; CAB09080.1; --
 CC EMBL: AF041819; AAB96962.1; --
 CC Biotin biosynthesis; Iron-sulfur; Transferase.
 FT METAL 85 85 IRON-SULFUR (POTENTIAL).
 FT METAL 89 89 IRON-SULFUR (POTENTIAL).
 FT METAL 92 92 IRON-SULFUR (POTENTIAL).
 SQ SEQUENCE 349 AA; 37550 MW; 3180B9DA CRC32;

 Query Match 57.4%; Score 31; DB 1; Length 349;
 Best Local Similarity 50.0%; Pred. No. 19;
 Matches 6; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
 QY 1 YRLAIXRIALRY 12
 | | | | |
 Db 280 FRALPRTMLRF 291

 RESULT 6
 SPAS_SALTY STANDARD; PRT; 356 AA.
 ID SPAS_SALTY
 AC P40702;
 DT 01-FEB-1995 (Rel. 31, Created)
 DT 01-FEB-1995 (Rel. 31, Last sequence update)
 DT 01-NOV-1995 (Rel. 32, Last annotation update)
 DE SURFACE PRESENTATION OF ANTIGENS PROTEIN SPAS.
 GN SPAS.
 OS Salmonella typhimurium.
 OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 OC Salmonella.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE: 94008985.
 RA GROISMAN E.A., OCHMAN H.;
 RT "Cognate gene clusters govern invasion of host epithelial cells by
 RL Salmonella typhimurium and Shigella flexneri."
 RL EMBO J. 12:3779-3787(1993).
 CC -1- FUNCTION: INVOLVED IN A SECRETORY PATHWAY RESPONSIBLE FOR THE
 CC SURFACE PRESENTATION OF DETERMINANTS NEEDED FOR THE ENTRY OF
 CC SALMONELLA SPECIES INTO MAMMALIAN CELLS.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. INNER MEMBRANE
 CC (POTENTIAL).
 CC -1- SIMILARITY: BELONGS TO THE FLHB/HRPN/YSCU/SPAS FAMILY.
 CC -----
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 CC -----

DR EMBL; X73525; CAA51927.1; -
 DR PIR; S37356; S37356
 DR STYGENE; SG10471; SPAS.
 DR PFAM; PF01312; Bac_export_2; 1.
 KW Virulence; Transmembrane; Inner membrane.
 FT TRANSMEM 29 49 POTENTIAL.
 FT TRANSMEM 72 92 POTENTIAL.
 FT TRANSMEM 132 152 POTENTIAL.
 FT TRANSMEM 179 199 POTENTIAL.
 FT TRANSMEM 261 281 POTENTIAL.
 SQ SEQUENCE 336 AA; 40093 MW; 26881BE1 CRC32;

Query Match 57.4%; Score 31; DB 1; Length 356;
 Best Local Similarity 50.0%; Pred. No. 19;
 Matches 6; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 YRLAIXRLRY 12
 I I I I I
 Db 70 YSLAVGIGLKY 81

RESULT 7

YD45_SCHPO STANDARD; PRT; 456 AA.
 AC Q10299;
 DT 01-OCT-1996 (Rel. 34, Created)
 DT 01-OCT-1996 (Rel. 34, Last sequence update)
 DT 01-OCT-1996 (Rel. 34, Last annotation update)
 DE HYPOTHETICAL 50.5 KD PROTEIN C22H10.05C IN CHROMOSOME I.
 GN SPAC22H10.05C.
 OS Schizosaccharomyces pombe (Fission yeast).
 OC Eukaryota; Fungi; Ascomycota; Archiascomycetes;
 OC Schizosaccharomycetales; Schizosaccharomycetaceae;
 CC Schizosaccharomycetes.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-972;
 RA DEVLIN K., CHURCHER C.M., BARRELL B.G., RAJANDREAM M.A., WALSH S.V.;
 RL Submitted (FEB-1996) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: TO C.ELEGANS F59A2.4.
 CC -----

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 CC -----

DR EMBL; Z69730; CAA93606.1; -
 KW Hypothetical protein; ATP-binding
 FT NP_BIND 113 120 ATP (POTENTIAL).
 SQ SEQUENCE 456 AA; 50449 MW; 11934D38 CRC32;

Query Match 57.4%; Score 31; DB 1; Length 456;
 Best Local Similarity 50.0%; Pred. No. 25;
 Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 1 YRLAIXRLAL 10
 I I I I I
 Db 205 YKLSLSRLAL 214

RESULT 8

PRPR_SALTY STANDARD; PRT; 541 AA.
 AC P74839;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 01-NOV-1997 (Rel. 35, Last annotation update)
 DE PROPIONATE CATABOLISM OPERON REGULATORY PROTEIN.
 GN PRPR.
 OS Salmonella typhimurium.
 OC Bacteria; Proteobacteria; gamma subdivision; Enterobacteriaceae;
 CC Salmonella.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-LT2;
 RX MEDLINE; 97158691.
 RA HORSWILL A.R., ESCALANTE-SEMERENA J.C.;
 FT "Propionate catabolism in Salmonella typhimurium LT2: two divergently
 RT transcribed units comprise the prp locus at 8.5 centisomes, prpR
 RT encodes a member of the sigma-54 family of activators, and the
 RT prpBCDE genes constitute an operon."
 RL J. Bacteriol. 179:928-940(1997).

CC -1- FUNCTION: INVOLVED IN THE TRANSCRIPTIONAL REGULATION OF THE
 CC PROPIONATE CATABOLISM OPERON.
 CC -1- SIMILARITY: THE CENTRAL REGION CONTAINS A SIGMA-54 FACTOR
 CC INTERACTION ATP-BINDING DOMAIN.
 CC -----

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 CC -----

DR EMBL; U51879; AAC44813.1; -
 DR STYGENE; SG22222; PRPR.
 DR PROSITE; PS00675; SIGMA54_INTERACT_1; FALSE_NEG.
 DR PROSITE; PS00676; SIGMA54_INTERACT_2; 1.
 DR PROSITE; PS00688; SIGMA54_INTERACT_3; 1.
 DR PROSITE; PS00689; SIGMA54_INTERACT_4; 1.
 DR PFAM; PF00158; sigma54; 1.
 KW Transcription regulation; DNA-binding; ATP-binding.
 FT DOMAIN 221 464 SIGMA-54 FACTOR INTERACTION (POTENTIAL).
 FT NP_BIND 321 330 ATP (POTENTIAL).
 FT DNA_BIND 513 532 H-T-H MOTIF (BY SIMILARITY).
 SQ SEQUENCE 541 AA; 60338 MW; 298D809F CRC32;

Query Match 57.4%; Score 31; DB 1; Length 541;
 Best Local Similarity 60.0%; Pred. No. 30;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 1 YRLAIXRLAL 10
 I I I I I
 Db 389 YRLSILRLTL 398

RESULT 9

YTFM_HAEIN STANDARD; PRT; 578 AA.
 AC P44038;
 DT 01-NOV-1995 (Rel. 32, Created)
 DT 01-NOV-1995 (Rel. 32, Last sequence update)
 DT 01-NOV-1995 (Rel. 32, Last annotation update)
 DE HYPOTHETICAL PROTEIN HI0698 PRECURSOR.
 GN HI0698
 OS Haemophilus influenzae.
 OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
 CC Haemophilus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-RD / KW20;
 RX MEDLINE; 95350630.

RA FLEISCHMANN R.D., ADAMS M.D., WHITE O., CLAYTON R.A., KIRKNESS E.F.,
 RA KERLAVAGE A.R., BULT C.J., TOMB J.-F., DOUGHERTY B.A., MERRICK J.M.,
 RA MCKENNEY K., SUTTON G., FITZHUGH W., FIELDS C.A., GOCAYNE J.D.,
 RA SCOTT J.D., SHIRLEY R., LIU L.-I., GLODE A., KELLEY J.M.,
 RA WEIDMAN J.F., PHILLIPS C.A., SPRIGGS T., HEDBLOM E., COTTON M.D.,
 RA UTERBACK T.R., HANNA M.C., NGUYEN D.T., SAUDEK D.M., BRANDON R.C.,
 RA FINE L.D., FRITCHMAN J.D., FUHRMANN J.L., GEORGE N.S.M.,
 RA GNEHM C.L., McDONALD L.A., SMALL K.V., FRASER C.M., SMITH H.O.,
 RA VENTER J.C.;
 RT "Whole-genome random sequencing and assembly of Haemophilus
 RL influenzae Rd.";
 RL Science 269:496-512(1995).
 CC -!- SIMILARITY: STRONG, TO E.COLI YTFM.
 CC -----
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 CC -----
 CC EMBL; U32752; AAC22357.1; -;
 DR TIGR; HI0698; -;
 KW Hypothetical protein; Signal.
 FT SIGNAL 1 22 POTENTIAL.
 FT CHAIN 23 578 HYPOTHETICAL PROTEIN HI0698.
 SQ SEQUENCE 578 AA; 65812 MW; 8FCD6AF9 CRC32;

Query Match 57.4%; Score 31; DB 1; Length 578;

Best Local Similarity 60.0%; Pred. NO. 32;

Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLAIXRLAL 10

I::|I::|I

Db 147 YKTAISRLAL 156

RESULT 10

MNT_HUMAN

ID MNT_HUMAN STANDARD; PRT; 582 AA.

AC Q99583;

DT 15-DEC-1998 (Rel. 37, Created)

DT 15-DEC-1998 (Rel. 37, Last sequence update)

DT 15-DEC-1998 (Rel. 37, Last annotation update)

DE MAX BINDING PROTEIN MNT (ROX PROTEIN) (MYC ANTAGONIST MNT).

GN MNT OR ROX.

OS Homo sapiens (Human).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

OC Eutheria; Primates; Catarrhini; Homnidae; Homo.

RN [1]

RP SEQUENCE FROM N.A.

RC TISSUE=FEETAL BRAIN;

RX MEDLINE; 97327566.

RA MERONI G., REYMOND A., ALCALAY M., BORSANI G., TANIGAMI A.,

RA TONLORENZI R., LO NIGRO C., MESSALI S., ZOLLO M., LEDBETTER D.H.,

RA BRENT R., BALLABIO A., CARROZZO R.;

RT "Rox, a novel bHLHZip protein expressed in quiescent cells that

RT heterodimerizes with Max, binds a non-canonical E box and acts as a

RT transcriptional repressor.";

RL EMBO J. 16:2892-2906(1997).

RN [2]

RP SEQUENCE FROM N.A.

RX MEDLINE; 98260677.

RA NIGRO C.L., VENESIO T., REYMOND A., MERONI G., ALBERICI P.,

RA CALINCARA S., ENRICO F., STACK M., LEDBETTER D.H., LISCIA D.S.,

RA BALLABIO A., CARROZZO R.;

RT "The human ROX Gene: genomic structure and mutation analysis in human

RT breast tumors.";

RL Genomics 49:275-282(1998).

CC -!- FUNCTION: BINDS DNA AS A HETERODIMER WITH MAX AND REPRESSES

CC TRANSCRIPTION. BINDS TO THE CANONICAL E BOX SEQUENCE

CC 5'-CACGTG-3' AND, WITH HIGHER AFFINITY, TO 5'-CACCG-3'.
 CC -!- SUBUNIT: EFFICIENT DNA BINDING REQUIRES DIMERIZATION WITH ANOTHER
 CC BHLH PROTEIN. BINDS DNA AS AN HOMODIMER OR A HETERODIMER WITH MAX.
 CC -!- SUBCELLULAR LOCATION: NUCLEAR.
 CC -!- SIMILARITY: BELONGS TO THE BASIC HELIX-LOOP-HELIX (BHLH) FAMILY OF
 CC TRANSCRIPTION FACTORS. BHLH-ZIP SUBFAMILY.
 CC -----

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CC EMBL; X96401; CAA65265.1; -;

DR EMBL; Y13440; CAA73851.1; -;

DR EMBL; Y13441; CAA73851.1; JOINED.

DR EMBL; Y13442; CAA73851.1; JOINED.

DR EMBL; Y13443; CAA73851.1; JOINED.

DR EMBL; Y13444; CAA73851.1; JOINED.

DR MIM; 603039; -;

DR PROSITE; PS00038; HELIX_LOOP_HELIX; FALSE_NEG.

DR PFAM; PF00010; HLH; 1.

KW Transcription regulation; Repressor; Nuclear protein; DNA-binding.

FT DNA_BIND 222 233 BASIC MOTIF (POTENTIAL).

FT DNA_BIND 234 270 HELIX-LOOP-HELIX MOTIF (POTENTIAL).

FT DOMAIN 271 299 LEUCINE-ZIPPER.

SQ SEQUENCE 582 AA; 62299 MW; 4ED96BBF CRC32;

Query Match 57.4%; Score 31; DB 1; Length 582;

Best Local Similarity 60.0%; Pred. NO. 32;

Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 3 LAIXRLALRY 12

I::|I::|I

Db 258 LSVLRALRY 267

RESULT 11

MNT_MOUSE

ID MNT_MOUSE STANDARD; PRT; 591 AA.

AC O08789; P97349;

DT 15-DEC-1998 (Rel. 37, Created)

DT 15-DEC-1998 (Rel. 37, Last sequence update)

DT 15-DEC-1998 (Rel. 37, Last annotation update)

DE MAX BINDING PROTEIN MNT (ROX PROTEIN) (MYC ANTAGONIST MNT).

GN MNT OR ROX.

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

RN [1]

RP SEQUENCE FROM N.A.

RC TISSUE=EMBRYO;

RX MEDLINE; 97152466.

RA HURLIN P.J., QUEVA C., EISENMAN R.N.;

RT "Mnt, a novel Max-interacting protein is coexpressed with Myc in

RT proliferating cells and mediates repression at Myc binding sites.";

RL Genes Dev. 11:44-58(1997).

RN [2]

RP SEQUENCE FROM N.A.

RC TISSUE=EMBRYO;

RX MEDLINE; 97327566.

RA MERONI G., REYMOND A., ALCALAY M., BORSANI G., TANIGAMI A.,

RA TONLORENZI R., LO NIGRO C., MESSALI S., ZOLLO M., LEDBETTER D.H.,

RA BRENT R., BALLABIO A., CARROZZO R.;

RT "Rox, a novel bHLHZip protein expressed in quiescent cells that

RT heterodimerizes with Max, binds a non-canonical E box and acts as a

RT transcriptional repressor.";

RL EMBO J. 16:2892-2906(1997).

CC -!- FUNCTION: BINDS DNA AS A HETERODIMER WITH MAX AND REPRESSES

CC TRANSCRIPTION. BINDS TO THE CANONICAL E BOX SEQUENCE 5'-CACGTG-3'

AND, WITH HIGHER AFFINITY, TO 5'-CAGCG-3'.

-1- SUBUNIT: EFFICIENT DNA BINDING REQUIRES DIMERIZATION WITH ANOTHER BHLH PROTEIN. BINDS DNA AS AN HOMODIMER OR A HETERODIMER WITH MAX. SUBCELLULAR LOCATION: NUCLEAR.

-1- SIMILARITY: BELONGS TO THE BASIC HELIX-LOOP-HELIX (BHLH) FAMILY OF TRANSCRIPTION FACTORS. BHLH-2IP SUBFAMILY.

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EMBL: U77356; AAB38687.1; -
 EMBL: Y07609; CAA68878.1; -
 MGD: MGI:109150; MNT.
 DR PROSITE; PS00038; HELIX_LOOP_HELIX; FALSE_NEG.
 DR PFAM; PF00010; HLH; 1.
 KW Transcription regulation; Repressor; Nuclear protein; DNA-binding.
 FT DNA_BIND 224 235 BASIC MOTIF (POTENTIAL).
 FT DOMAIN 236 272 HELIX-LOOP-HELIX MOTIF (BY SIMILARITY).
 FT DOMAIN 273 301 LEUCINE-ZIPPER (POTENTIAL).
 FT CONFLICT 379 392 T -> P (IN REF. 2).
 FT CONFLICT 392 392 V -> A (IN REF. 2).
 FT CONFLICT 402 403 EE -> QQ (IN REF. 2).
 FT CONFLICT 414 414 G -> A (IN REF. 2).
 FT CONFLICT 431 431 V -> A (IN REF. 2).
 FT CONFLICT 465 465 A -> P (IN REF. 2).
 FT CONFLICT 525 525 T -> A (IN REF. 2).
 FT CONFLICT 558 558 G -> A (IN REF. 2).
 SQ SEQUENCE 591 AA; 63311 MW; 16BADBA6 CRC32;

Query Match 57.48; Score 31; DB 1; Length 591;
 Best Local Similarity 60.08; Pred. No. 33;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 3 LAIXRIALRY 12
 I:: I I I I I
 Db 260 LSVLTALRY 269

RESULT 12

ID YPD2 CAEEL STANDARD; PRT; 833 AA.
 AC Q11182;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 01-NOV-1997 (Rel. 35, Last annotation update)
 DE HYPOTHETICAL 93.8 KD PROTEIN C05D11.2 IN CHROMOSOME III.
 GN C05D11.2
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RC SEQUENCE FROM N.A.
 RP STRAIN-BRISTOL N2;
 DU Z;
 RA Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
 CC -1- SIMILARITY: WEAK, TO YEAST VPS16.
 CC -----
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EMBL: U00048; AAB53823.1; -
 DR WORMPEP; C05D11.2; CE01132.

KW Hypothetical protein.
 SQ SEQUENCE 833 AA; 93791 MW; 508777F2 CRC32;

Query Match 57.48; Score 31; DB 1; Length 833;
 Best Local Similarity 70.08; Pred. No. 48;
 Matches 7; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

Qy 1 YRLAIXRIAL 10
 I I I I I I
 Db 589 YELAISRIPL 598

RESULT 13

ID CYA6_MOUSE STANDARD; PRT; 1165 AA.
 AC Q01341;
 DT 01-APR-1993 (Rel. 25, Created)
 DT 01-APR-1993 (Rel. 25, Last sequence update)
 DT 15-DEC-1999 (Rel. 39, Last annotation update)
 DE ADENYLATE CYCLASE, TYPE VI (EC 4.6.1.1) (ATP PYROPHOSPHATE-LYASE) (CA(2+)-INHIBITABLE ADENYLYL CYCLASE).
 GN ADCY6
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 92357702.
 RA YOSHIMURA M., COOPER D.M.F.;
 RT "Cloning and expression of a Ca(2+)-inhibitable adenylyl cyclase from NCB-20 cells.";
 RT Proc. Natl. Acad. Sci. U.S.A. 89:6716-6720(1992).
 RN [2]
 RP SEQUENCE OF 10-1365 FROM N.A.
 RX MEDLINE; 93076707.
 RA FREMONT R.T., JACOBOWITZ O., IYENGAR R.;
 RT "Lowered responsiveness of the catalyst of adenylyl cyclase to stimulation by GS in heterologous desensitization: a role for adenosine 3',5'-monophosphate-dependent phosphorylation.";
 RT Endocrinology 131:2774-2784(1992).
 CC -1- FUNCTION: THIS A MEMBRANE-BOUND, CA(2+)-INHIBITABLE ADENYLYL CYCLASE.
 CC -1- CATALYTIC ACTIVITY: ATP = 3',5'-CYCLIC AMP + PYROPHOSPHATE.
 CC -1- ENZYME REGULATION: INHIBITION BY CA(2+) IN THE SUBMICROMOLAR CONCENTRATION RANGE.
 CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
 CC -1- TISSUE SPECIFICITY: MOST ABUNDANT IN HEART BUT WEAKLY DETECTABLE IN BRAIN, INTESTINE, LUNG, AND SPLEEN.
 CC -1- DOMAIN: COMPOSED OF TWO HOMOLOGOUS DOMAINS.
 CC -1- SIMILARITY: BELONGS TO ADENYLYL CYCLASE CLASS-4/GUANYLYL CYCLASE FAMILY.
 CC -----
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EMBL: M93422; AAA37174.1; -
 EMBL: M96653; AAA37182.1; -
 DR PIR; A46187; A46187.
 DR HSSP; P19754; LAWK.
 DR MGD; MGI:87917; ADCY6
 DR PROSITE; PS00452; GUANYLATE_CYCLASES; 2.
 DR PFAM; PF00211; guanylate_cyc; 2.
 KW Lyase; GMP synthetase; Transmembrane; Glycoprotein; Duplication.
 FT DOMAIN 1 149 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 150 166 POTENTIAL.
 FT TRANSMEM 179 195 POTENTIAL.
 FT TRANSMEM 212 228 POTENTIAL.

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FT TRANSMEM 237 253 POTENTIAL.
FT TRANSMEM 257 273 POTENTIAL.
FT TRANSMEM 287 303 POTENTIAL.
FT DOMAIN 304 670 CYTOPLASMIC (POTENTIAL).
FT TRANSMEM 671 688 POTENTIAL.
FT TRANSMEM 699 715 POTENTIAL.
FT TRANSMEM 740 756 POTENTIAL.
FT DOMAIN 757 816 EXTRACELLULAR (POTENTIAL).
FT TRANSMEM 817 833 POTENTIAL.
FT TRANSMEM 836 852 POTENTIAL.
FT TRANSMEM 894 910 POTENTIAL.
FT DOMAIN 911 1165 CYTOPLASMIC (POTENTIAL).
FT CARBOHYD 277 277 POTENTIAL.
FT CARBOHYD 790 790 POTENTIAL.
FT CARBOHYD 875 875 POTENTIAL.
FT CONFLICT 47 47 K -> N (IN REF. 2).
FT CONFLICT 76 76 G -> A (IN REF. 2).
FT CONFLICT 508 509 GR -> RAG (IN REF. 2).
FT CONFLICT 737 737 V -> G (IN REF. 2).
FT CONFLICT 881 881 L -> Q (IN REF. 2).
FT CONFLICT 990 990 V -> M (IN REF. 2).
SQ SEQUENCE 1165 AA; 130318 MW; 485F7155 CRC32;

Query Match 57.4%; Score 31; DB 1; Length 1165;
Best Local Similarity 55.6%; Pred. No. 69;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 4 AIXRIALRY 12
I:|:|:|
Db 885 AVGRVALKY 893

RESULT 14
CYA6_RAT
ID QYA6_RAT STANDARD; PRT; 1166 AA.
AC Q03343;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE ADENYLATE CYCLASE, TYPE VI (EC 4.6.1.1) (ATP PYROPHOSPHATE-LYASE)
DE CA(2+)-INHIBITABLE ADENYLYL CYCLASE).
GN ACY6.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 93077589.
RA KRUPINSKI J., LEHMAN T.C., FRANKENFIELD C.D., ZWAAGSTRA J.C.,
RA WATSON P.A.;
RT "Molecular diversity in the adenylylcyclase family. Evidence for
RT eight forms of the enzyme and cloning of type VI."
RL J. Biol. Chem. 267:24858-24862(1992).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=LIVER;
RX MEDLINE; 93028552.
RA PREMONT R.T., CHEN J., MA H.-W., PONNAPALLI M., IYENGAR R.;
RT "Two members of a widely expressed subfamily of hormone-stimulated
RT adenylyl cyclases."
RL Proc. Natl. Acad. Sci. U.S.A. 89:9809-9813(1992).
CC -1- FUNCTION: THIS A MEMBRANE-BOUND, CA(2+)-INHIBITABLE ADENYLYL
CC CYCLASE.
CC -1- CATALYTIC ACTIVITY: ATP = 3',5'-CYCLIC AMP + PYROPHOSPHATE.
CC -1- ENZYME REGULATION: INHIBITION BY CA(2+) IN THE SUBMICROMOLAR
CC CONCENTRATION RANGE.
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -1- DOMAIN: COMPOSED OF TWO HOMOLOGOUS DOMAINS.
CC -1- SIMILARITY: BELONGS TO ADENYLYL CYCLASE CLASS-4/GUANYLYL CYCLASE
CC FAMILY.
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CC or send an email to license@isb-sib.ch).
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CC EMBL; L01115; AAA40676.1; -
CC EMBL; M96160; AAA40678.1; ALT_INIT.
CC PIR; A45145; A45145.
CC HSP; P19754; LMK.
CC PROSITE; PS00452; GUANYLATE_CYCLASES; 2.
CC PFAM; PF00211; guanylate_cyc; 2.
CC Lysase; CAMP synthesis; Transmembrane; Glycoprotein; Duplication.
CC DOMAIN 1 149 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 150 166 POTENTIAL.
CC TRANSMEM 179 195 POTENTIAL.
CC TRANSMEM 212 228 POTENTIAL.
CC TRANSMEM 237 253 POTENTIAL.
CC TRANSMEM 257 273 POTENTIAL.
CC TRANSMEM 287 303 POTENTIAL.
CC DOMAIN 304 671 CYTOPLASMIC (POTENTIAL).
CC TRANSMEM 672 689 POTENTIAL.
CC TRANSMEM 700 716 POTENTIAL.
CC TRANSMEM 741 757 POTENTIAL.
CC DOMAIN 758 817 EXTRACELLULAR (POTENTIAL).
CC TRANSMEM 818 834 POTENTIAL.
CC TRANSMEM 837 853 POTENTIAL.
CC TRANSMEM 895 911 POTENTIAL.
CC DOMAIN 912 1166 CYTOPLASMIC (POTENTIAL).
CC CARBOHYD 791 791 POTENTIAL.
CC CARBOHYD 876 876 POTENTIAL.
CC CONFLICT 80 80 K -> E (IN REF. 2).
CC CONFLICT 130 130 R -> P (IN REF. 2).
CC CONFLICT 538 538 G -> A (IN REF. 2).
CC CONFLICT 790 790 I -> L (IN REF. 2).
SQ SEQUENCE 1166 AA; 130506 MW; AA449BC2 CRC32;

Query Match 57.4%; Score 31; DB 1; Length 1166;
Best Local Similarity 55.6%; Pred. No. 69;
Matches 5; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 4 AIXRIALRY 12
I:|:|:|
Db 886 AVGRVALKY 894

RESULT 15
LMA3_MOUSE
ID LMA3_MOUSE STANDARD; PRT; 2569 AA.
AC Q61789; Q61788; Q61966;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE LAMININ ALPHA-3 CHAIN PRECURSOR (FRAGMENT).
GN LMA3.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-BALB/C; TISSUE=LUNG;
RX MEDLINE; 95394948.
RA GALLIANO M.-F., ABERDAM D., AGUZZI A., ORTONNE J.-P., MENEGUZZI G.;
RT "Cloning and complete primary structure of the mouse laminin alpha 3
RT chain. Distinct expression pattern of the laminin alpha 3A and alpha
RL J. Biol. Chem. 270:21820-21826(1995).
RN [2]
RP REVISIONS.
RA ABERDAM D.;
RL Submitted (APR-1997) to the EMBL/GenBank/DBJ databases.

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NON_TER	1	28	29	2569	3 CHAIN.	POTENTIAL.
SIGNAL	<1	28	29	2569	LAMININ ALPHA-3 CHAIN.	
CHAIN	29	2569	29	2569	DOMAIN IV'.	
DOMAIN	29	498	29	498	DOMAIN III B.	
DOMAIN	499	700	499	700	3.5 X LAMININ EGF-LIKE REPEATS.	
DOMAIN	546	700	546	700	LAMININ EGF-LIKE 1.	
DOMAIN	546	589	546	589	LAMININ EGF-LIKE 2.	
DOMAIN	590	639	590	639	LAMININ EGF-LIKE 3.	
DOMAIN	640	690	640	690	LAMININ EGF-LIKE 4 (N-TERMINAL).	
DOMAIN	691	700	691	700	LAMININ DOMAIN IV (DOMAIN IV A).	
DOMAIN	701	889	701	889	3 X LAMININ EGF-LIKE REPEATS (DOMAIN III A).	
DOMAIN	890	1057	890	1057	LAMININ EGF-LIKE 4 (C-TERMINAL).	
DOMAIN	890	922	890	922	LAMININ EGF-LIKE 5.	
DOMAIN	923	959	923	959	LAMININ EGF-LIKE 6.	
DOMAIN	970	1022	970	1022	LAMININ EGF-LIKE 7 (INCOMPLETE).	
DOMAIN	1023	1057	1023	1057	DOMAIN II AND I (HEPTAT REPEATS).	
DOMAIN	1058	1648	1058	1648	5 X LAMININ G-LIKE REPEATS (DOMAIN G).	
DOMAIN	1649	2569	1649	2569	LAMININ G-LIKE 1.	
DOMAIN	1649	1825	1649	1825	LAMININ G-LIKE 2.	
DOMAIN	1826	1934	1826	1934	LAMININ G-LIKE 3.	
DOMAIN	1995	2209	1995	2209	LAMININ G-LIKE 4.	
DOMAIN	2210	2385	2210	2385	LAMININ G-LIKE 5.	
DOMAIN	2386	2569	2386	2569	COILED COIL (POTENTIAL).	
DOMAIN	1090	1219	1090	1219	COILED COIL (POTENTIAL).	
DOMAIN	1251	1296	1251	1296	COILED COIL (POTENTIAL).	
DOMAIN	1327	1404	1327	1404	COILED COIL (POTENTIAL).	
DOMAIN	1450	1477	1450	1477	COILED COIL (POTENTIAL).	
DOMAIN	1557	1622	1557	1622	CELL ATTACHMENT SITE (POTENTIAL).	
SITE	1513	1515	1513	1515	BY SIMILARITY.	
DISULFID	546	553	546	553	BY SIMILARITY.	
DISULFID	548	560	548	560	BY SIMILARITY.	
DISULFID	562	571	562	571	BY SIMILARITY.	
DISULFID	574	587	574	587	BY SIMILARITY.	
DISULFID	590	605	590	605	BY SIMILARITY.	
DISULFID	592	612	592	612	BY SIMILARITY.	
DISULFID	614	623	614	623	BY SIMILARITY.	
DISULFID	626	637	626	637	BY SIMILARITY.	
DISULFID	640	652	640	652	BY SIMILARITY.	
DISULFID	642	659	642	659	BY SIMILARITY.	
DISULFID	661	670	661	670	BY SIMILARITY.	
DISULFID	673	688	673	688	BY SIMILARITY.	
DISULFID	923	932	923	932	BY SIMILARITY.	
DISULFID	925	939	925	939	BY SIMILARITY.	
DISULFID	942	951	942	951	BY SIMILARITY.	
DISULFID	954	967	954	967	BY SIMILARITY.	
DISULFID	970	982	970	982	BY SIMILARITY.	
DISULFID	972	991	972	991	BY SIMILARITY.	
DISULFID	993	1002	993	1002	BY SIMILARITY.	
DISULFID	1005	1020	1005	1020	INTERCHAIN (PROBABLE).	
DISULFID	1058	1058	1058	1058	INTERCHAIN (PROBABLE).	
DISULFID	1061	1061	1061	1061	POTENTIAL.	
CARBOHYD	591	591	591	591	POTENTIAL.	
CARBOHYD	912	912	912	912	POTENTIAL.	
CARBOHYD	1398	1398	1398	1398	POTENTIAL.	
CARBOHYD	1500	1500	1500	1500	POTENTIAL.	
CARBOHYD	1571	1571	1571	1571	POTENTIAL.	
CARBOHYD	1600	1600	1600	1600	POTENTIAL.	
CARBOHYD	1737	1737	1737	1737	POTENTIAL.	
CARBOHYD	1819	1819	1819	1819	POTENTIAL.	
CARBOHYD	1986	1986	1986	1986	POTENTIAL.	
CARBOHYD	2333	2333	2333	2333	POTENTIAL.	
CARBOHYD	2509	2509	2509	2509	POTENTIAL.	
VARSPIC	1	842	1	842	MISSING (IN ISOFORM A).	
VARSPIC	843	901	843	901	ATVLPRLRLHRLHFTETORLTGLVGLLEASDTGSGPR MHVEMCAQCPDVTGDS -> MLPVVRWSAWSTGWLWFG AALGCGYSGEQQRQVAFLOPQSNHLQASMYELRPS (IN ISOFORM A).	

SEQUENCE

Query Match 57.4%; Score 31; DB 1; Length 2569;
Best Local Similarity 60.0%; Pred. NO. 1.6e+02;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 3 LAIXRIALRY 12
i::|||
Db 15 LSLFRIVLY 24

Search completed: February 8, 2000, 00:59:58
Job time: 3787 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:46 ; Search time 209.03 seconds
(without alignments)
3.980 Million cell updates/sec

Title: US-08-653-294-19
Perfect score: 54
Sequence: 1 YRLAIXRLRY 12

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 225878 seqs, 69334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : SPTREMBL12.*
1: SP_archaea.*
2: SP_bacteria.*
3: SP_fungi.*
4: SP_human.*
5: SP_invertebrate.*
6: SP_mammal.*
7: SP_mhc.*
8: SP_organelle.*
9: SP_phage.*
10: SP_plant.*
11: SP_rodent.*
12: SP_virus.*
13: SP_vertebrate.*
14: SP_unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	38	70.4	325	2 Q9XB05	Q9XB05 myxococcus
2	35	64.8	795	5 Q22063	Q22063 caenorhabdi
3	35	64.8	1899	10 Q9XEG1	Q9XEG1 gossypium h
4	34	63.0	141	2 O84514	O84514 chlamydia t
5	34	63.0	309	2 O86347	O86347 mycobacteri
6	33	61.1	142	2 Q92759	Q92759 chlamydia p
7	32	59.3	264	2 O84015	O84015 chlamydia t
8	32	59.3	310	2 O86606	O86606 streptomyce
9	32	59.3	610	2 P73400	P73400 synecocyst
10	31	57.4	42	11 O54741	O54741 mus musculu
11	31	57.4	190	2 O67313	O67313 aquifex aeo
12	31	57.4	243	2 O85853	O85853 sphingomona
13	31	57.4	407	10 O23555	O23555 arabidopsis
14	31	57.4	469	1 O28589	O28589 archaeoglob
15	31	57.4	910	13 Q9YGE5	Q9YGE5 oncorhynch
16	31	57.4	983	3 O94545	O94545 schizosacch
17	31	57.4	1847	5 P91495	P91495 caenorhabdi
18	30	55.6	171	2 O69220	O69220 azotobacter
19	30	55.6	206	2 Q923X8	Q923X8 pseudomonas
20	30	55.6	211	2 P76343	P76343 escherichia

21	30	55.6	236	2	O88071	O88071 streptomyce
22	30	55.6	286	1	O29923	O29923 archaeoglob
23	30	55.6	313	2	P97213	P97213 clostridium
24	30	55.6	376	1	P95964	P95964 sulfolobus
25	30	55.6	377	2	O9WZD6	O9WZD6 thermotoga
26	30	55.6	394	2	Q47265	Q47265 escherichia
27	30	55.6	430	5	O4725	O4725 caenorhabdi
28	30	55.6	441	5	Q26954	Q26954 trypanosoma
29	30	55.6	497	2	P76767	P76767 escherichia
30	30	55.6	573	4	O9V4B9	O9V4B9 homo sapien
31	30	55.6	880	2	O3ZD55	O3ZD55 rickettsia
32	30	55.6	984	4	O43718	O43718 homo sapien
33	30	55.6	1084	2	O83423	O83423 treponema p
34	30	55.6	1147	5	O9XZ87	O9XZ87 drosophila
35	30	55.6	1604	10	O9X140	O9X140 arabidopsis
36	29	53.7	35	5	O61237	O61237 onchocerca
37	29	53.7	89	7	O19569	O19569 homo sapien
38	29	53.7	90	7	O46697	O46697 gorilla gor.
39	29	53.7	104	12	O11430	O11430 avian adeno
40	29	53.7	127	8	O31747	O31747 chara coral
41	29	53.7	133	7	O19189	O19189 homo sapien
42	29	53.7	133	12	O85384	O85384 variola vir
43	29	53.7	138	1	O26976	O26976 methanobact
44	29	53.7	138	7	O78209	O78209 homo sapien
45	29	53.7	144	1	Q919C4	Q919C4 aeropyrum p

ALIGNMENTS

RESULT 1

Q9XB05 PRELIMINARY; PRT; 325 AA.
AC Q9XB05;
DT 01-NOV-1999 (TREMBLrel. 12, Created)
DT 01-NOV-1999 (TREMBLrel. 12, Last sequence update)
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
DE MEMBRANE ASSOCIATED PROTEIN.
GN TAD.
OS Myxococcus xanthus.
OC Bacteria; Proteobacteria; delta subdivision: Myxobacteria;
OC Myxococcales; Cystobacterineae; Myxococcaceae; Myxococcus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-ER-15;
RA PAITAN Y., ORR E., RON E.Z., ROSENBERG E.;
RT "Genetic and functional analysis of genes required for the post-
RT modification of the polyketide antibiotic TA of Myxococcus xanthus.";
RL Submitted (JAN-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL; AJ132503; CAB46503.1; -
SQ SEQUENCE 325 AA; 35985 MW; 4CC64E85 CRC32;

Query Match 70.4%; Score 38; DB 2; Length 325;
Best Local Similarity 58.3%; Pred. No. 2.9;
Matches 7; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIXRLRY 12

DB 202 YRLTVDRPLRY 213

RESULT 2

Q22063 PRELIMINARY; PRT; 795 AA.
AC Q22063; Q93489;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-MAY-1999 (TREMBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
DE T01C3.10 PROTEIN.
GN T01C3.10
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;

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OC Rhabditiina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RA WILD A.;
RL Submitted (AUG-1996) to the EMBL/GenBank/DBJ databases.
DR EMBL; Z81061; CAB02939.1; -.
DR EMBL; Z78413; CAB02939.1; JOINED.
DR EMBL; Z78413; CAB01667.1; -.
DR EMBL; Z81061; CAB01667.1; JOINED.
DR HSP; P19491; IGR2.
DR PFAM; PF00060; l1g_chan; 1.
SQ SEQUENCE 795 AA; 89703 MW; DD722166 CRC32;

Query Match 64.8%; Score 35; DB 5; Length 795;
Best Local Similarity 50.0%; Pred. No. 31;
Matches 6; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIXRLRY 12
   |||: |||
Db 6 YRTSLRLATRY 17

RESULT 3
ID Q9XEG1 PRELIMINARY; PRT; 1899 AA.
AC Q9XEG1;
DT 01-NOV-1999 (TEMBLrel. 12, Created)
DT 01-NOV-1999 (TEMBLrel. 12, Last sequence update)
DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
DE PUTATIVE CALLOSE SYNTHASE CATALYTIC SUBUNIT.
GN CFL1.
OS Gossypium hirsutum (Upland cotton).
OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;
OC core eudicots; Rosidae; eurosids II; Malvales; Malvaceae; Gossypium.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CV. TEXAS MARKER-1; TISSUE=PRIMARY-STAGE COTTON FIBER;
RA CUI X., SHIN H., BROWN R.M.;
RT "Cotton CFL1 gene shows homology to the yeast beta-1,3-glucan synthase
RT subunit FKS1."
RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AF085717; AAD25952.1; -.
SQ SEQUENCE 1899 AA; 218627 MW; E695145F CRC32;

Query Match 64.8%; Score 35; DB 10; Length 1899;
Best Local Similarity 63.8%; Pred. No. 79;
Matches 7; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAXRLRY 12
   |||: |||
Db 554 RLAVSRFLRF 564

RESULT 4
ID O84514 PRELIMINARY; PRT; 141 AA.
AC O84514;
DT 01-NOV-1998 (TEMBLrel. 08, Created)
DT 01-NOV-1998 (TEMBLrel. 08, Last sequence update)
DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
DE 50S RIBOSOMAL PROTEIN L17.
GN RL17.
OS Chlamydia trachomatis.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=D/UN-3/CX;
RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L.,
RA MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V.,
RA DAVIS R.W.;

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RT "genome sequence of an obligate Intracellular Pathogen of Humans:
RT Chlamydia trachomatis.";
RN Science 0:0-0(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=D/UN-3/CX;
RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L.,
RA MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V.,
RA DAVIS R.W.;
RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: BELONGS TO THE L17P FAMILY OF RIBOSOMAL PROTEINS.
DR EMBL; AE001323; AAC68107.1; -.
DR PROSITE; PS01167; RIBOSOMAL_L17; 1.
DR PFAM; PF01196; Ribosomal_L17; 1.
KW Ribosomal protein.
SQ SEQUENCE 141 AA; 16152 MW; 2570FF7 CRC32;

Query Match 63.0%; Score 34; DB 2; Length 141;
Best Local Similarity 54.5%; Pred. No. 8.3;
Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAXRLRY 12
   |||: |||
Db 64 RLAVGRLMRY 74

RESULT 5
ID O86347 PRELIMINARY; PRT; 309 AA.
AC O86347;
DT 01-NOV-1998 (TEMBLrel. 08, Created)
DT 01-NOV-1998 (TEMBLrel. 08, Last sequence update)
DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
DE HYPOTHETICAL 33.5 KD PROTEIN.
GN RV276C.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE; 98295987.
RA COLE S.T., BROCH S., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
RA GORDON S.V., BIGLMEIER K., GAS S., BARRY III C.E., TEKAIA F.,
RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
RA DAVIES R., DEVLIN K., FELTWEILL T., GENTLES S., HAMLIN N., HOLROYD S.,
RA HORNSBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,
RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAM M.A., ROGERS J.,
RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence."
RL Nature 393:537-544(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RA PARKHILL J.;
RL Submitted (JUN-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AL008967; CAAL5591.1; -.
DR HSP; P33184; 2PIA.
DR PROSITE; PS00197; 2FE2S_FERREDOXIN; 1.
DR PFAM; PF00111; fer2; 1.
DR PFAM; PF00175; oxidore_fad; 1.
KW Hypothetical protein; Iron-sulfur.
SQ SEQUENCE 309 AA; 33517 MW; B152B590 CRC32;

Query Match 63.0%; Score 34; DB 2; Length 309;
Best Local Similarity 77.8%; Pred. No. 19;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIXRIA 9

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Db 65 YRIAIRRIA 73
|||||

RESULT 6
Q927S9 PRELIMINARY; PRT: 142 AA.
ID Q927S9
AC Q927S9
DT 01-MAY-1999 (TRENBLrel. 10, Created)
DT 01-MAY-1999 (TRENBLrel. 10, Last sequence update)
DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
DE 505 RIBOSOMAL PROTEIN L17.
GN RL17.
OS Chlamydia pneumoniae.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydophila.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CWL029;
RA KALMAN S., MITCHELL W., MARATHE R., LAMMEL C., FAN J., OLINGER L., GRIMWOOD J., DAVIS R.W., STEPHENS R.S.;
RT "Comparative Genomes of Chlamydia pneumoniae and C. trachomatis.";
RL Submitted (DEC-1998) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: BELONGS TO THE L17p FAMILY OF RIBOSOMAL PROTEINS.
DR EMBL: AE001647; AAD18764.1; -.
KW PROSITE; PS01167; RIBOSOMAL_L17; 1.
SQ SEQUENCE 142 AA; 16400 MW; 4839EC84 CRC32;

Query Match 61.1%; Score 33; DB 2; Length 142;
Best Local Similarity 54.5%; Pred. No. 14;
Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

Qy 2 RLAIKRIALRY 12
|:|:|:|:|
Db 64 RLAIKRLMVRY 74

RESULT 7
O84015 PRELIMINARY; PRT: 264 AA.
ID O84015
AC O84015
DT 01-NOV-1998 (TRENBLrel. 08, Created)
DT 01-NOV-1998 (TRENBLrel. 08, Last sequence update)
DT 01-NOV-1998 (TRENBLrel. 08, Last annotation update)
DE HYPOTHETICAL 30.0 KD PROTEIN.
GN YBBP.
OS Chlamydia trachomatis.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=D/UV-3/CX;
RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L., MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V., DAVIS R.W.;
RT "Genome Sequence of an Obligate Intracellular Pathogen of Humans: Chlamydia trachomatis.";
RL Science 0:0-0(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=D/UV-3/CX;
RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L., MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V., DAVIS R.W.;
RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL: AE001276; AAC67602.1; -.
KW Hypothetical protein.
SQ SEQUENCE 264 AA; 29971 MW; C0AD12DB CRC32;

Query Match 59.3%; Score 32; DB 2; Length 264;
Best Local Similarity 70.0%; Pred. No. 42;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

Qy 2 RLAIKRIALR 11
|:|:|:|:|
Db 83 RLALSRIALR 92

RESULT 8
O86606 PRELIMINARY; PRT: 310 AA.
ID O86606
AC O86606;
DT 01-NOV-1998 (TRENBLrel. 08, Created)
DT 01-NOV-1998 (TRENBLrel. 08, Last sequence update)
DT 01-NOV-1998 (TRENBLrel. 08, Last annotation update)
DE PUTATIVE TRANSPOSASE.
GN SC3A7.05C.
OS Streptomyces coelicolor.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Streptomycineae; Streptomycetaceae; Streptomyces.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=A3(2);
RA OLIVER K., HARRIS D.;
RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=A3(2);
RA PARKHILL J., BARRELL B.G., RAJANDREAM M.A.;
RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RC STRAIN=A3(2);
RX MEDLINE; 97000351.
RA REDENBACH M., KIESER H.M., DENAPATE D., EICHNER A., CULLUM J., KINASHI H., HOPWOOD D.A.;
RT "A set of ordered cosmids and a detailed genetic and physical map for the 8 Mb Streptomyces coelicolor A3(2) chromosome.";
RL Mol. Microbiol. 21:77-96(1996).
DR EMBL; AL031155; CAZ00868.1; -.
SQ SEQUENCE 310 AA; 33156 MW; 5AD2CED4 CRC32;

Query Match 59.3%; Score 32; DB 2; Length 310;
Best Local Similarity 63.6%; Pred. No. 50;
Matches 7; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

Qy 2 RLAIKRIALRY 12
|:|:|:|:|
Db 249 RLAVRWATRY 259

RESULT 9
P73400 PRELIMINARY; PRT: 610 AA.
ID P73400
AC P73400;
DT 01-FEB-1997 (TRENBLrel. 02, Created)
DT 01-FEB-1997 (TRENBLrel. 02, Last sequence update)
DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
DE ABC TRANSPORTER.
OS Synechocystis sp. (strain PCC 6803).
OC Bacteria; Cyanobacteria; Chroococcales; Synechocystis.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=PCC6803;
RA TABATA S.;
RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=PCC6803;
RX MEDLINE; 97061201.
RA KANEKO T., SATO S., KOTANI H., TANAKA A., ASAMIZU E., NAKAMURA Y., MIYAJIMA N., HIROSAWA M., SUGIURA M., SASAMOTO S., KIMURA T., HOSOUCHI T., MATSUNO A., MURAKI A., NAKAZAKI N., NARUO K., OKUMURA S., SHIMPO S., TAKEUCHI C., WADA T., WATANABE A., YAMADA M.,

RA TABATA S.;
 RT "Sequence analysis of the genome of the unicellular cyanobacterium
 RT Synechocystis sp. strain PCC6803. II. Sequence determination of the
 RT entire genome and assignment of potential protein-coding regions."
 RL DNA Res 3:109-136 (1996).
 DR ENBL; D90906; BAA17440.1; -.
 DR HSP; P13569; INBD.
 DR PFAM; PF00664; ABC_membrane; 1.
 DR PFAM; PF00005; ABC_tran; 1.
 SQ SEQUENCE 610 AA; 67028 MW; 835D2045 CRC32;

Query Match 59.38; Score 32; DB 2; Length 610;
 Best Local Similarity 70.08; Pred. No. 1e+02;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIKXRIALR 11
 ||||| |||
 Db 508 RLAIARAWR 517

RESULT 10
 O54741
 ID O54741 PRELIMINARY; PRT; 42 AA.
 AC O54741;
 DT 01-JUN-1998 (TrEMBLrel. 06, Created)
 DT 01-JUN-1998 (TrEMBLrel. 06, Last sequence update)
 DT 01-NOV-1999 (TrEMBLrel. 12, Last annotation update)
 DE LAMININ, ALPHA 3 (LAMININ 5 ALPHA3C CHAIN) (FRAGMENT).
 GN LAMA3.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN-SV129;
 RC MEDLINE; 97400537.
 RA FERRIGNO O., VIROLLE T., GALLIANO M.F., CHAUVIN N., ORTONNE J.P.,
 RA MENEGUZZI G., ABERDAM D.;
 RT "Murine laminin alpha3A and alpha3B isoform chains are generated by
 RT usage of two promoters and alternative splicing."
 RL J. Biol. Chem. 272:20502-20508 (1997).
 DR ENBL; Y08850; CAA70073.1; -.
 DR MGD; MGI:99909; Lama3.
 FT NON_TER 42
 SQ SEQUENCE 42 AA; 4872 MW; 42FE6150 CRC32;

Query Match 57.48; Score 31; DB 11; Length 42;
 Best Local Similarity 60.08; Pred. No. 10;
 Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 3 LAIXRIALRY 12
 ||: |||
 Db 22 LSLFRIVLY 31

RESULT 11
 O67313
 ID O67313 PRELIMINARY; PRT; 190 AA.
 AC O67313;
 DT 01-AUG-1998 (TrEMBLrel. 07, Created)
 DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
 DT 01-NOV-1998 (TrEMBLrel. 08, Last annotation update)
 DE HYPOTHETICAL 22.0 KD PROTEIN.
 GN AQ_1277.
 OS Aquifex aeolicus.
 OC Bacteria; Aquificales; Aquificaceae; Aquifex.
 [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN-VF5;
 RX MEDLINE; 98196665.
 RA DECKERT G., WARREN P.V., GAASTERLAND T., YOUNG W.G., LENOX A.L.,

RA GRAHAM D.E., OVERBECK R., SNEAD M.A., KELLER M., AUJAY M., HUBER R.,
 RA FELDMAN R.A., SHORT J.M., OLSON G.J., SWANSON R.V.;
 RT "The complete genome of the hyperthermophilic bacterium Aquifex
 RT aeolicus."
 RL Nature 392:353-358 (1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-VF5;
 RA DECKERT G., WARREN P.V., GAASTERLAND T., YOUNG W.G., LENOX A.L.,
 RA GRAHAM D.E., OVERBECK R., SNEAD M.A., KELLER M., AUJAY M., HUBER R.,
 RA FELDMAN R.A., SHORT J.M., OLSON G.J., SWANSON R.V.;
 RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
 DR ENBL; AE00732; AAC07278.1; -.
 KW Hypothetical protein.
 SQ SEQUENCE 190 AA; 22041 MW; EFB61F50 CRC32;

Query Match 57.48; Score 31; DB 2; Length 190;
 Best Local Similarity 54.58; Pred. No. 49;
 Matches 6; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLAIXRIALR 11
 : ||| :|||
 Db 43 HNLAIQKVALR 53

RESULT 12
 O85853
 ID O85853 PRELIMINARY; PRT; 243 AA.
 AC O85853;
 DT 01-NOV-1998 (TrEMBLrel. 08, Created)
 DT 01-NOV-1998 (TrEMBLrel. 08, Last sequence update)
 DT 01-NOV-1998 (TrEMBLrel. 08, Last annotation update)
 DE HYPOTHETICAL 26.5 KD PROTEIN.
 OS Sphingomonas aromaticivorans.
 OC Plasmid pNL1.
 OC Bacteria; Proteobacteria; alpha subdivision; Sphingomonas group;
 OC Sphingomonas.
 [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN-F199;
 RA ROMINE M.F., STILLWELL L.C., WONG K.-K., THURSTON S.J., SISK E.C.,
 RA SENSEN C.W., GAASTERLAND T., SAFFER J.D., FREDRICKSON J.K.;
 RT "Complete sequence of a 184 kb catabolic plasmid from Sphingomonas
 RT aromaticivorans strain F199."
 RL Submitted (JUL-1998) to the EMBL/GenBank/DBJ databases.
 DR ENBL; AF079317; AAD03868.1; -.
 KW Hypothetical protein; Plasmid.
 SQ SEQUENCE 243 AA; 26455 MW; 40CDFBF4 CRC32;

Query Match 57.48; Score 31; DB 2; Length 243;
 Best Local Similarity 54.58; Pred. No. 63;
 Matches 6; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 RLAIKXRIALRY 12
 ||| | :|||
 Db 5 RLAIQRRVTIRY 15

RESULT 13
 O23555
 ID O23555 PRELIMINARY; PRT; 407 AA.
 AC O23555;
 DT 01-JAN-1998 (TrEMBLrel. 05, Created)
 DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TrEMBLrel. 12, Last annotation update)
 DE PUTATIVE SERINE PROTEASE-LIKE PROTEIN.
 GN DL4585C.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons;
 OC core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;

OC Arabidopsis.

RN [1]
 RP SEQUENCE FROM N.A.
 RA BEVAN M., STEKEMA W., MURPHY G., WAMBUIT R., POHL T., TERRY N.,
 RA KREIS M., KAVANAGH T., ENTIAN K.D., RIEGER M., JAMES R.,
 RA PUGDOMENEC P., HATZIOPOULOS P., OBERMAIER B., DUESTERHOFF A.,
 RA JONES J., PALME K., ANSGORGE W., DELSENY M., BANCROFT I., MEWES H.W.,
 RA SCHUELLER C., CHALWATZIS N.;
 RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA EU ARABIDOPSIS SEQUENCING PROJECT;
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 DR EMBL; 297342; CAB46052.1; -;
 KW Protease.
 SQ SEQUENCE 407 AA; 46703 MW; 9B2D1556 CRC32;

Query Match 57.4%; Score 31; DB 10; Length 407;

Best Local Similarity 58.3%; Pred. No. 1.1e+02;

Matches 7; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

Qy 1 YRLAIXRIALRY 12

Db 368 YRLGNRIALSF 379

RESULT 14

O28589
 ID O28589 PRELIMINARY; PRT; 469 AA.
 AC O28589;
 DT 01-JAN-1998 (TRENBLrel. 05, Created)
 DT 01-JAN-1998 (TRENBLrel. 05, Last sequence update)
 DT 01-AUG-1998 (TRENBLrel. 07, Last annotation update)
 DE CONSERVED HYPOTHETICAL PROTEIN.
 GN AF1684.
 OS Archaeoglobus fulgidus.
 OC Archaea; Euryarchaeota; Archaeoglobales; Archaeoglobaceae;
 OC Archaeoglobus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-VC-16 / DSM 4304 / ATCC 49558;
 RX MEDLINE; 98049343.
 RA KLENK H.-P., CLAYTON R.A., TOMB J.-F., WHITE O., NELSON K.E.,
 RA KETCHUM K.A., DODSON R.J., GWINN M., HICKEY E.K., PETERSON J.D.,
 RA RICHARDSON D.L., KERLAVAGE A.R., GRAHAM D.E., KYRIDES N.C.,
 RA FLEISCHMANN R.D., QUACKENBUSH J., LEE N.H., SUTTON G.G., GILL S.,
 RA KIRKNESS E.F., DOUGHERTY B.A., MCKENNEY K., ADAMS M.D., LOFTUS B.,
 RA PETERSON S., REICH C.I., MCNEIL L.K., BADGER J.H., GLODER A., ZHOU L.,
 RA OVERBECK R., GOCAYNE J.D., WEIDMAN J.F., McDONALD L., UTTERBACK T.,
 RA COTTON M.D., SPRIGGS T., ARTIACH P., KAINE B.P., SYKES S.M.,
 RA SADOW P.W., D'ANDREA K.P., BOWMAN C., FUJII C., GARLAND S.A.,
 RA MASON T.M., OLSEN G.J., FRASER C.M., SMITH H.O., WOESE C.R.,
 RA VENTER J.C.;
 RT "The complete genome sequence of the hyperthermophilic, sulphate-
 reducing archaeon Archaeoglobus fulgidus."
 RL Nature 390:364-370(1997).
 DR EMBL; AE000987; AAB89565.1; -;
 DR TIGR; AF1684; -;
 KW Hypothetical protein.
 SQ SEQUENCE 469 AA; 53468 MW; 9D430846 CRC32;

Query Match 57.4%; Score 31; DB 1; Length 469;

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Db 12 YRKGKRIALRY 23

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 DT 01-MAY-1999 (TRENBLrel. 10, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
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 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Actinopterygii;
 OC Neopterygii; Teleostei; Euteleostei; Protacanthopterygii;
 OC Salmoniformes; Salmonidae; Oncorhynchus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 99132290.
 RA BRUNELLI J.P., THORGAARD G.H.;
 RT "Sequence, expression and genetic mapping of a rainbow trout
 retinoblastoma cDNA";
 RL Gene 226:175-180(1999).
 DR EMBL; AF102861; AAD13390.1; -;
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57.4%; Score 31; DB 13; Length 910;

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Search completed: February 8, 2000, 13:17:48

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Blazej, R.G., Chavez, C., Chew, M., Doyle, C.M., Farfan, D.E.,
Flanagan, J., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L.,
Kim, S.H., Lee, B., Lomontan, M.A., Mak, J., Mazda, P., Moshrefi, A.R.,
Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S., Pfeiffer, B., Punch, E.,
Snir, E., Twomey, B., Wan, K.H., Zhang, R., Zieran, L.L. and Rubin, G.M.
Sequencing of Drosophila chromosome 3L, region 61F3-62A2
Unpublished (1997)

2 (bases 1 to 268369)

Celniker, S.E., George, R.A., Galle, R.F., Hoskins, R.A.,
Swirskes, R.R., Harris, N.L., Agapayni, A., Arcaina, T.T., Baxter, E.,
Blazej, R.G., Chavez, C., Chew, M., Doyle, C.M., Farfan, D.E.,
Flanagan, J., Houston, K.A., Hummasti, S.R., Karra, K., Kearney, L.,
Kim, S.H., Lee, B., Lomontan, M.A., Mak, J., Mazda, P., Moshrefi, A.R.,
Moshrefi, M., Nixon, K., Pacleb, J.M., Park, S., Pfeiffer, B., Punch, E.,
Snir, E., Twomey, B., Wan, K.H., Zhang, R., Zieran, L.L. and Rubin, G.M.
Direct Submission
Submitted (22-OCT-1998) Berkeley Drosophila Genome Project, MS
64-121, Lawrence Berkeley National Laboratory, One Cyclotron Road,
Berkeley, CA 94720, US
Sequence submitted by:
Berkeley Drosophila Genome Project
Lawrence Berkeley National Laboratory, MS 64-121
Berkeley, CA 94720
For further information about this sequence, including its location
and relationship to other sequences, please visit our sequence
archive Web site (<http://www.fruitfly.org/sequence/>) or send email
to bdgpp@fruitfly.berkeley.edu.
Library locations: 11-34, 21-67, 44-36, 73-50, 92-76.

Location/Qualifiers
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Protein, *tac* gene, *tac* gene, *tac* gene, *tac* gene, *tac* gene:
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Myxococcus xanthus.
Myxococcus xanthus.
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Bacteria; Proteobacteria; della subdivISION; Mycobacteria;

Myxococcales; Cystobacterineae; Myxococcaceae; Myxococcus.
 1 (bases 1 to 4513)
 Paitan, Y., Orr, E., Ron, E.Z., and Rosenberg, E.
 Genetic and functional analysis of genes required for the
 post-modification of the polyketide antibiotic TA of Myxococcus
 xanthus
 Unpublished
 2 (bases 1 to 4513)
 Paitan, Y.
 Direct Submission
 Submitted (25-JAN-1999) Paitan Y., Molecular Microbiology and
 Biotechnology, Tel Aviv University, G.S. Wise Faculty of Life
 Sciences, Tel Aviv University, Ramat Aviv, 69978, ISRAEL
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 VERSION U72788.1 GI:1575796
 KEYWORDS HTG.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE
 1 (bases 1 to 25383)
 Sulston, J.E. and Waterston, R.
 TITLE Toward a complete human genome sequence
 JOURNAL Genome Res. 8 (11), 1097-1108 (1998)
 MEDLINE 99063792
 REFERENCE 2 (bases 1 to 25383)
 Waterston, R.
 TITLE The sequence of Homo sapiens cosmid clone U169D2
 JOURNAL Unpublished (1999)
 REFERENCE 3 (bases 1 to 25383)
 Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (29-SEP-1996)
 REFERENCE 4 (bases 1 to 25383)
 Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (19-JAN-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 REFERENCE 5 (bases 1 to 25383)
 Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (20-JAN-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 REFERENCE 6 (bases 1 to 25383)
 Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (20-JAN-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 REFERENCE 7 (bases 1 to 25383)
 Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (20-JAN-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 REFERENCE 8 (bases 1 to 25383)
 Waterston, R.
 TITLE Direct Submission
 JOURNAL Submitted (20-JAN-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

TITLE Direct Submission
 JOURNAL Submitted (27-Apr-1999) Department of Genetics, Washington
 University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
 COMMENT SUBMITTED BY: WUGSC
 Genome Sequencing Center
 Department of Genetics
 Washington University
 St. Louis MO 63108, USA
 http://genome.wustl.edu/gsc
 mailto:sapiens@watson.wustl.edu

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
 all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

This clone was mapped by Grieff, M., Whyte, M. P., Thakker, R. V., and Mazzarella, R. Sequence analysis of 139 kb in Xp22.1 containing spermine synthase and the 5' region of PEX. Genomics 44:227-231 (1997).

SOURCE INFORMATION:

This clone is from a chromosome X-specific cosmid library LLOXNCC01 'U'. The source of the chromosomes was a human/hamster hybrid, GM07297-F, from Robert Nussbaum at the University of Pennsylvania School of Medicine. Please contact the Lawrence Livermore National Laboratory at <http://www-bio.llnl.gov/genome> to obtain the clone.

FEATURES

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AUTHORS	2 (bases 1 to 36947)		hemolytica glycoproteinase Al and E.coli yj3C);
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AUTHORS	Robison,K.		/transl_table=11
TITLE	Direct Submission		/product="u229e"
JOURNAL	Submitted (01-NOV-1993) Department of Genetics, Harvard Medical		/protein_id="AAAL7310.1"
	School, 200 Longwood Avenue, Boston MA 02115		/db_xref="GI:467128"
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AUTHORS	Robison,K.		VDEQARFGVPEIASRAHLEALGPTIRCALAAAGLTGSAKPDVVAATIGPGLAGALL
JOURNAL	Submitted (01-MAR-1994) Department of Genetics, Harvard Medical		VGVAARAYSAAGVPFYAVNHLGHLAADVHEGPLPCVALLVSGGTHLQVRSLL
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JOURNAL	located at Collaborative Research Incorporated (1365 Main St.,		LIVGVGAANSRCAS"
	Waltham MA, 02159). 617-487-7979). Please contact Doug Smith		complement(5386..6465)
COMMENT	(smith@cr.cric.com). The annotation should be considered		/EC_number="2.3.1.128"
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	LLHOKISITPELLPELVKTESGKPLLIIVADELGEALATLVNIRKTLKAVAVKS		complement(7931..9097)
	PPFDRRKAFLEDLAIVTGGVYVNPETGLVREVGTDVLGSRARVYVSKDDTIIVDGG		/EC_number="5.1.1.1"
	GSDNAVKRYNQLRAIEVSDSEWDRKQLQERVAKLAGGVAVIKGVAVTETALKRKE		/note="Alanine racemase; B229_C3_243"
	SVEDAVAAAKASIEEGIIAGGSGALVQCGAALKQLTSLTGDEALGIDVFEALKAPL		/codon_start=1
	YWTATNAGLGVAVVVDKVSGLPAGHGLNASTLGLYGLVADGVDPVKVTRSAVLNAA		/transl_table=11
			/product="alr"
			/protein_id="AAAL7309.1"

seq_documentation_block:
LOCUS MLCB2533 40245 bp DNA BCT 27-AUG-1999
DEFINITION Mycobacterium leprae cosmid B2533.
ACCESSION AL035310
VERSION AL035310.1 GI:4200258
KEYWORDS ansp; ATP-dependent RNA helicase; ATP-phosphoribosyl transferase; hisG 5-methyltetrahydrofolate-homocysteine methyl transferase; hisI; L-asparagine permease; mtbB;

```

/note="MLCB2533.01c, ansp, probable L-asparagine permease,
partial CDS, len: >366 aa; highly similar to many
amino-acid permeases e.g. ANSP_SALTY (EMBL:U04851)
S.typhimurium Ansp, L-asparagine permease (L-asparagine
transport protein) (497 aa), fasta scores; opt: 1508
z-score: 1696.8 E(): 0, 61.2% identity in 366 aa overlap.
Equivalent to M.tuberculosis Rv2127, ansp. (MTCY261.26,
85.7% identity in 356 aa overlap). Also similar to
M.tuberculosis permease Rv0346c, aroP2 (MTCY13E10.06c,
75.1% identity in 345 aa overlap). Probable integral
membrane protein, contains PS00218 Amino acid permeases
signature. Pfam match to entry PF00324 aa_permeases, amino
acid permease. Annotated as ORF TR:Q49801, designated
aroP2 in M.leprae cosmid EMBL:U00017"
/codon_start=1
/transl_table=11
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/protein_id="CAA22915.1"
/db_xref="GI:4200259"
/translation="MATLAESPCKSGASRAGVIGEGYHKGLKPRQLQMIGIGGAI
GTGLFCAGRLAKAGPGLFVYAVCGVFILRALGELVLRHPSGGSFVSAREFF
GEKAAVVGWLYFLDWMATIAVDTTATVTLHRTTFTALPQWTLALLAVLVNML
ISVWEGELFEWALIKVCALMAFLVVGIFLGGRYPDGHTGLSLWTSGLGLEPTG
VAQLIVSSGWFAYAAVELVGTAGETVEPKIMPRAINSVIARAIYVGSVILLA
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INGSGPKFTARMSKNGVPYGGIILAAVICLG"
1..1168
/note="1168 bp perfect direct repeat"
/complement(2..991)
/gene="ansp"
/note="Pfam match to entry PF00324 aa_permeases, Amino
acid permease, score 245.80, E-value 6e-70"
438
/unsure

/note="ansp"
/note="ambiguous base T /G"
/complement(833..925)
/gene="ansp"
/note="PS00218 Amino acid permeases signature"
1223
/note="conflict: T is TG EMBL:ML017"
/complement(1249..2766)
/gene="ansp2"
/complement(1249..2766)
/note="ansp2"
/note="MLCB2533.02c, ansp2, probable L-asparagine
permease, len: 505 aa; highly similar to many amino-acid
permeases e.g. ANSP_SALTY (EMBL:U04851) S.typhimurium
Ansp, L-asparagine permease (L-asparagine transport
protein) (497 aa), fasta scores;opt: 1891 z-score: 2218.9
E(): 0, 58.9% identity in 477 aa overlap. Equivalent to
M.tuberculosis Rv2127 (MTCY261.26, 83.7% identity in 485
aa overlap). Also similar to M.tuberculosis permease
Rv0346c (MTCY13E10.06c, 69.8% identity in 473 aa overlap).
Probable integral membrane protein, contains PS00218 Amino
acid permeases signature. Pfam match to entry PF00324
aa_permeases, amino acid permease. Annotated as ORF
TR:Q49802, designated lysP in M.leprae cosmid EMBL:U00017"
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GTGLFCAGRLAKAGPGLFVYAVCGVFILRALGELVLRHPSGGSFVSAREFF
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ISVWEGELFEWALIKVCALMAFLVVGIFLGGRYPDGHTGLSLWTSGLGLEPTG
VAQLIVSSGWFAYAAVELVGTAGETVEPKIMPRAINSVIARAIYVGSVILLA
LLPYSAFKASESPFTFFSKVGFYAGDLMNIVLTAASSLNAGLYATGRVWMSIA
WGTIVLCQLRHKRMKIMRRFRMRPLAPYSGYLTLAFILAVLVMAFDKPIGTWT
VASLIVTPALIAWYTSIRKRVNTIARRMVGTGPPPAIANPVPQPSERSHQNP"
complement(1354..2691)
/gene="ansp2"

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/note="Pfam match to entry PF00324 aa_permeases, Amino
acid permease, score 501.80, E-value 5.1e-147"
1668..2835
/note="1168 bp perfect direct repeat"
2104
/gene="ansp2"
/note="conflict: C is CT in EMBL:ML017"
/complement(2500..2592)
/gene="ansp2"
/note="PS00218 Amino acid permeases signature"
/complement(2903..3856)
/gene="MLCB2533.03c"
/complement(2903..3856)
/gene="MLCB2533.03c"
/note="MLCB2533.03c, hypothetical protein, len: 317 aa;
similar to M.tuberculosis hypothetical protein Rv2125
(MTCY261.21) (EMBL:Z97559) (292 aa), fasta scores; opt:
1848 z-score: 2382.2 E(): 0, 84.1% identity in 290 aa
overlap. Also some similarity to M.leprae hypothetical
protein TR:Q49847 (29.7% identity in 279 aa overlap).
Annotated as ORF TR:Q49797, hypothetical protein in
M.leprae cosmid EMBL:U00017"
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/db_xref="GI:4200261"
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/translation="MPPHRAVTRHASSALKPYADSVTLRDGPDGALPELHNTVVV
AFEGWNSDASGALEHLNVAWEADPVEIDDEAYDYQVNRVPIQVDGVITRELY
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TPHTRPVPSGAAYSPESARREGLEETRYEGTIGAGVQDACVAARIIPAVMFMAAVP
HYVSHPPKPAVKATVALLRVEDVLDVEVLADLPQAEDEWQAITEAEDDELAETVH
SLEQRGAEDVDNALGKIDGDAAEFERYLRRRRPGFGR"
4004..7555
/gene="meth"
4004..7555
/gene="meth"
/note="MLCB2533.04, meth, probable
5-methyltetrahydrofolate-homocysteine methyltransferase,
len: 1183 aa; similar to many members of vitamin-B12
dependent methionine synthase family e.g. METH_ECOLI
(EMBL:X15584) E.coli meth (1226 aa), fasta scores; opt:
1617 z-score: 1000.7 E(): 0, 31.6% identity in 1228 aa
overlap. Equivalent to M.tuberculosis Rv2124c
(MTCY261.20c, 88.7% identity in 1183 aa overlap).
Annotated as METH_MYCLE, designated meth2 in M.leprae

```

```

alignment_scores:
Quality: 38.00 Length: 12
Ratio: 4.222 Gaps: 0
Percent Similarity: 75.000 Percent Identity: 66.667

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alignment_block:
US-08-653-294-19 x MLCB2533 ..

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Align seg 1/1 to: MLCB2533 from: 1 to: 40245

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1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||
17534 TATCGATGGCAATCGAGNATATCTGCACGGTAT 17569

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seq_name: gb_bai:U00017

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seq documentation_block:
LOCUS U00017 42157 bp DNA BCT 01-MAR-1994

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DEFINITION Mycobacterium leprae cosmid B2126.
ACCESSION U00017
VERSION U00017.1 GI:466994
KEYWORDS
SOURCE

```

```

Mycobacterium leprae.
Mycobacterium leprae
Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;

```

Actinomycetales: Corynebacterineae; Mycobacteriaceae;

REFERENCE

1 (bases 1 to 42157)

Smith, D.R.

Unpublished

2 (bases 1 to 42157)

Robison, K.

Direct Submission

Submitted (01-NOV-1993) Department of Genetics, Harvard Medical

School, 200 Longwood Avenue, Boston MA 02115

3 (bases 1 to 42157)

Robison, K.

Direct Submission

Submitted (01-MAR-1994) Department of Genetics, Harvard Medical

School, 200 Longwood Avenue, Boston MA 02115

On Mar 31, 1994 this sequence version replaced gi:414223.

This sequence data was produced by the Genome Sequencing Center

located at Collaborative Research Incorporated (1365 Main St.,

Waltham MA, 02159). 617-487-7979). Please contact Doug Smith

(smith@cr.cric.com). The annotation should be considered

preliminary and incomplete.

Location/Qualifiers

FEATURES

source

1. 42157

/organism="Mycobacterium leprae"

/db_xref="taxon:1769"

complement(133..1086)

/note="match to yigv and yigv E.coli; B2126_C1_183"

/codon_start=1

/transl_table=11

/product="u2126a"

/protein_id="AA17191.1"

/db_xref="GI:467006"

/translation="MNCRAVVRACDILLKRIKHYSRSTNPATMSLDHLTELRTF

LLISLAIVTTIFGFYWSHSIFGLESGLWLRPPYCLPQSRADISDPGQCLLA

TAFDQPMRLKIVGMAIGIVLSPVWFYQLWAFITPGLYTKERRTFVAFVPAVLFA

GGTVLVLKALGFLIIVSGVQVTFALSGDRYFELLNLLVVGVSFFEPILLVNL

NIAGLLTYORLKSRRGLIFAMFVFAVFTPGSDPFSMTALGAALTLLLELAQLVRL

HDKRRVKEALADDEASVIEPPSSIPERSYATRSDDVT"

complement(1132..1398)

/note="match to yigt E.coli; B2126_C1_182"

/codon_start=1

/transl_table=11

/product="u2126b"

/protein_id="AA17190.1"

/db_xref="GI:467005"

/translation="MGLSPHWVVLVVVVVLFAGAKKLPDAARSLGKSMRIFKSELR

EMOTENQAQASALETPNQNTVVSQSRVPPWSTEQDTEARPA"

complement(1471..1692)

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/transl_table=11

/product="B2126_C1_181"

/protein_id="AA17189.1"

/db_xref="GI:467004"

/translation="MSWIFVYPMQAHLPDGSFQAVMTYASDMMTRLVLGFSNV

QVQAPALAYVRNNAVALESQVTAQA"

complement(1701..2444)

/codon_start=1

/transl_table=11

/product="B2126_C2_220"

/protein_id="AA17199.1"

/db_xref="GI:467014"

/translation="MQLSRLVRLNKNVYFOANPRITRAEAAADLVSAKQLDQDF

NQWVGLPGYGGDLIDFEFGDTIEVTSAGIDRLPQLTSPKAGLLVALRALANI

PGVVDPEASAKTEAAADAVVMGNEATGVSVDTPPFSESHAVRAAVRAVRKQAL

VIDYYSASHDTLTSRIVDPTRVLLVGHSHLEAWSREAQGVRLFRFDRIVVARELDEP

AAAPETVRRRCQTHRSSMTTRCCRRRCG"

complement(2441..3436)

/codon_start=1

/transl_table=11

/product="B2126_C3_266"

/protein_id="AA17208.1"

/db_xref="GI:467023"

/translation="MAISKVERLVNLVIALSTLRGYMTAEKIRSSVAGYSDSPTEAF

SRMFERDKNELRLGLIPLEVGVKYSALDPSSEGYRINRDAYALPPVELTDPDAAYAVAT
QLWSEOLITATQGALKKRAAGVDIDPLDTPVVIASSSGVSSLRGSEDFLSILLSAI
GSRQAVQFPYRPAEPTYNVPEWGVITENSCWLVGHDCDRNATRTFRLSRIGSE
VAPIGAGAVTPDGVDLRRIVSDVAEAVSTGATARVWVVDGRATLRHAGRPAGVR
LGGRDGOVIELDIGSIDRLARDIAGHCADAVVLEPDALRDDVLRLRAHAGTGPS"

CDS

/codon_start=1

/transl_table=11

/product="B2126_C3_265"

/protein_id="AA17207.1"

/db_xref="GI:467022"

/translation="MTEIDWVLRKLFQYQDRDNMELTDPKIAQLDLAYHDKRGR

GVFDLQKGLAARYTDEDIADNHPPTTRANLRGEFISAAQAAGRFTVDWVHL

KLNDQAQRTVLCKDPPFRAVDKVRLLIAM"

complement(3861..4871)

/codon_start=1

/transl_table=11

/product="B2126_C2_219"

/protein_id="AA17198.1"

/db_xref="GI:467013"

/translation="MVGWYPAWLHRAHQTTSTLKDQVRRIMGIETFGVCTTFHGH

RLSPDEVARYLFRFVWGRSSNVFLRNGARLYLDVSHPEYATACDNLVLQVTHD

RAGEVLEDLDAEORLADEGGDIYLFKNNTDSAGNSYGCHEYLIVRAGEFSRI

SDVLEPLVTRQLICGAGKVLQPKAATFCLSORAHHWEGSVSSATRSPLNTRDE

PHADAERYRLHVIQSDNMCETTMLKVTALMLEMVEVTPGFDFDSLDNFIAR

EVSHDITGRPVRLAGRSALDIOREYITRAFEHLQTRPNVQFEQVVDLMGPSAR

RR"

complement(6569..6862)

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/product="B2126_C3_261"

/protein_id="AA17206.1"

/db_xref="GI:467021"

/translation="MKIADANVLLYSGTSSHTGHPCAGSTVRLSCADRIFGWVPL

LTVRPATKMGVLRTMSSDAIGQVADMLTGPSAVLMCLTVRHAFLVKILV"

complement(6822..7115)

/codon_start=1

/transl_table=11

/product="B2126_C2_217"

/protein_id="AA17197.1"

/db_xref="GI:467012"

/translation="MSFKKVFNDVIRDSVLGRFEPHFRTACITDLALSAVNLQWALPI

AAEFDEELVRLSGFGRHSRFLSHRLAPHRLPARIITHENRRERAVER"

complement(7253..8050)

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/transl_table=11

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/protein_id="AA17205.1"

/db_xref="GI:467020"

/translation="MSFPFISPEQAMRQRELARKIARGSVVALAYAGGVLFVAE

NPSRLQKISELYDRVGFAGAGFNFNLRGGIQFADTRGYAYDRDVTGRLANY

YAQTGTFTEQAKPYEVELCAVEAHYGETKPPELYRITYDGSINDEPHFWMGGTT

ESTANLKESYAENSLDALGIAVAALRAGSADAAGSDOPTLGVASLEVALDANRP

RRAFRIIGSGLEALLREKDKSGKAQNPKGARDSKSKSGESTD"

complement(8047..8922)

/note="proteaseome, beta subunit; B2126_C1_173"

/codon_start=1

/transl_table=11

/product="prCB"

/protein_id="AA17188.1"

/db_xref="GI:467003"

/translation="WTRSPDRLPNTLAFPGISVINQSFVDLLRQAPPELLPVSLGG

GOSGGQQSHGTTIVLKYPGGVYAGDRRSQGNRIAGDRVKVYIYDDYTATGIA

GIAVAVERFARLYAVELEHYKLEGVPLTFAGKVNRLAINVRNSNLAAQGLALPL

AGVDIHAPQPSAGRIIVSDAGGWNIEEGYQSGVSGSIFAKSSKIKLYSOVSADS

ALRVATEALYDAADDDSATGGPDLVRGIPTATVIGAEAGAEVETESRLAREITES

RSRATILSGFGSEK"

complement(9142..10686)

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/transl_table=11

/product="B2126_C1_172"

/protein_id="AA17187.1"

/db_xref="GI:467002"


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  /codon_start=1
  /protein_id="AAC67317.1"
  /db_xref="GI:3785970"
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AKTEPOKLOVDSKDYSRNTMLYLAAAGFLGNGFLNGEAAEAARVKRNRKKALEK
LRAKASEPNKSGNOKIEKELEKEVFLLPPLVVEANLQ"
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  complement(<1769..>.2932)
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  /note="hypothetical protein"
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  /db_xref="GI:3785971"
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RSPGLYPEGLTWPDQHLVGLSHSRTIHSVDAGVETLISDLDPENSTILGLAV
DSNRRLLACIQSLPLPFPFSAALASYDRSGRRVFLSPPLSLPDDDEDIAROVANDY
AVDEGNAYVTSNAKFIKWDGDAASIRSKPLNSQPAADADASFDCGLNGIV
YISGILLVQSTGKVKFVDEDSGNARLVLLGDLAAGDMTRRRDGTVMVVSQKK
LWLLKSDSSEGVSEYDEIDLDIEGFPFVAVTRDRIYVLYGRVMEGINSKKEG
ARWFGLEEVSEKGEEDIKWLYVLGFGFAYFCFVRFQMKLITNMKKIT"
  complement(3756..3826)
  /note="exon predicted by xgrail, quality good"
  complement(4609..4669)
  /note="exon predicted by xgrail, quality marginal"
  complement(join(6667..7104,7209..7285,7380..7537,
7915..8000,8140..8407,8565..>.9759))
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  complement(join(7038..7104,7209..7285,7380..7537,
7915..8000,8140..8407,8565..>.9759))
  /gene="F219.4"
  /codon_start=1
  /product="putative auxin transport protein"
  /protein_id="AAC67319.1"
  /db_xref="GI:3785972"
  /translation="MITWHDLYVLRVPLVYVAMILAYGVQVQWKKIFSPDQCSGINR
FVAIFPLPSFHFISINDPYAMNFREVAADTLQKIIMLVLLALWNLKNGSLEWMI
TIFSLPTLNTLVGMVPIPLIAMFTYAGSLMVQVVIQCIITWTLILFLFYRGAKLL
IMQFPTGASIVSFYKVESDVSLDGHDFLEDAEIGNCGKLVHVKRKSASRLMM
TPRPSNLTAIEIYLSLSTPRGSNFHSDFYVMVGPGRSLNFGPDLYSVQSRGPT
PRPSFNENAVYGFYNTNNSVPASPDYAPNPFSTGTGTVSTKPNKIPKQOOL
QKDSKASHDAKELHMFVWSSSPVSDVFGGAGDNVATEQSEQGAKEIRMVVSDOP
RKNAGGGDIDGLDSEGERIEKTAGLNKNGSNSTAELEAAGDGGNGTHMP
PTSVMTLIIIMVYKLRINPTNYSSLIIGLWALVYRWHVAMPKILQOQSILSDAG
LGMAMFSLGFLMALQPKIACGNSVATFAMAVRFTGPATMAVAGIAGLHLLRIA
IVQALPQGVPEVFAKEYNVHTLSTGVIFGMLIALPTLYIYILLGL"
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  10586..10649
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marginal_shadowexon"

10803..10939
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excellent_shadowexon"
  complement(11141..11230)
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  11620..11654
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marginal_shadowexon"
  complement(12452..12519)
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marginal_shadowexon"
  complement(13191..13230)
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marginal_shadowexon"
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NHTLNPKQKVEFLAKHLPRQIEVFNRRARSKLKQTEMECEYLKRWFGSLTEENH
RLRHEVEELRAMKVGTPTVNSASLSLTMPCRCERTVPAASPSRAVVPVPAKTFPPQER
DR"
  14881..14924
  /rpt_family="AT-rich"
  complement(16632..16710)
  /note="exon predicted by xgrail, quality excellent"
  complement(16998..17033)

alignment_scores:
  Quality: 38.00 Length: 11
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  Percent Similarity: 100.000 Percent identity: 72.727

alignment_block:
  US-08-653-294-19 x ATAC005560/rev ..
  Align seg 1/1 to reverse of: ATAC005560 from: 1 to: 103125

2 ArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
  ::::::::::::::::::::::::::::::::::::
  44994 AAGTTGGGAATTACTAGAGTTCATTAGATAT 44962

seq_name: gb_htg1:CNS01DTM
seq_documentation_block:
  LOCUS CNS01DTM 191442 bp DNA HTG 04-NOV-1999
  DEFINITION Homo sapiens chromosome 14 clone R-80A15, *** SEQUENCING IN
  PROGRESS ***, in ordered pieces.
  ACCESSION AL132800
  VERSION AL132800.1 GI:6272127
  KEYWORDS HTG; HTGS_PHASE2.
  SOURCE human.
  ORGANISM Homo sapiens
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
  Eutheria; Primates; Catarrhini; Hominidae; Homo.

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1 TyrArgLeuAlaIle***ArgIleAlaLeuArg 11
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 1727 TTTCGATGTCATCGCAGCATATCATTACGC 1695

seq_name: gb_pr3:HS782L23

seq_documentation_block:

LOCUS HS782L23 123925 bp DNA PRI 23-NOV-1999

DEFINITION Human DNA sequence from clone 782L23 on chromosome 1p31.2-33

Contains start of HOOK1 gene, ESTs, STS and GSSs, complete

sequence.

ACCESSION AL035416

VERSION AL035416.7 GI:4775629

KEYWORDS HTG: CPG Island.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;

Eutheria; Primates; Catarrhini; Hominidae; Homo.

1 (bases 1 to 123925)

Wallis.J.

Direct Submission

Submitted (04-AUG-1999) Sanger Centre, Hinxton, Cambridgeshire,

CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

requests: clonerequest@sanger.ac.uk

On May 11, 1999 this sequence version replaced gi:4741514.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

The following abbreviations are used to associate primary accession

numbers given in the feature table with their source databases:

Em: EMBL; Sw: SWISSPROT; Tr: TREMBL

This sequence is the entire insert of clone 782L23. This sequence

has been finished according to sequence map criteria as follows. An

attempt is made to resolve all sequencing problems, such as

compressions and repeats, but not necessarily within known

annotated human repeat sequence elements (e.g. Alu). Where the

sequence is ambiguous, there is an annotation using the 'unsure'

feature key.

This sequence was generated from part of bacterial clone contigs of

human chromosome 1, constructed by the Sanger Centre Chromosome 1

Mapping Group. Further information can be found at

<http://www.sanger.ac.uk/HGP/Chr1>

782L23 is from the library RPC14 constructed at the Roswell Park

Cancer Institute by the group of Pieter de Jong. For further

details see <http://bacpac.med.buffalo.edu/VECTOR:pcypac2>.

Location/Qualifiers

FEATURES
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1..123925

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/db_xref="taxon:9606"

/chromosome="1"

/map="p31.2-33"

/clone="RP4-782L23"

/clone_lib="RPC1-4"

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6225..6285,6755..6842,16087..16197,16754..16955,

18085..18225,21672..21838,24879..24962,28335..28397,

29858..29925,31733..31865,33057..33107,36418..36490,

43327..43412,50090..50318))

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/note="match: cDNAs: Em:AF044923; match: ESTs:

Em:AA161507"

/evidence=not_experimental

/product="dj782L23.1 (HOOK1)"

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/gene="dj782L23.1"

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18085..18225,21672..21838,24879..24962,28335..28397,

29858..29925,31733..31865,33057..33107,36418..36490,

43327..43412,50090..50152))
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 SKELSSPPDAVGELEQLKRALEELQALAEKELRQCEELDMQVTTIQDENSL
 VSENMENKLDQDSDPNTVAKKIFHAQLOLEQLQENFLEAKDDIRVHCE
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 DLKQVKTQETNMVHTVSLLEELKANAARQLETYRQVODLHVKLSSSEKRA
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 134..253
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 823..880
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 1468..1513
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 2074..2177
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 3183..3362
 /note="3 copies 60 mer 96% conserved"
 3533..3766
 /note="MIR repeat: matches 7..237 of consensus"
 3846..3966
 /note="MIR repeat: matches 3..139 of consensus"
 4059..4194
 /note="AluJb repeat: matches 1..142 of consensus"
 4195..4506
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 4507..4684
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 6554..6726
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 7728..9025
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 9026..9279
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 9281..9320
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 9321..9497
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 9532..9611
 /note="L1M4 repeat: matches 1466..1553 of consensus"
 9645..10125
 /note="L1P1A1 repeat: matches 5665..6148 of consensus"
 10164..10429
 /note="L1M4 repeat: matches 987..1206 of consensus"
 10441..10920
 /note="L1MB7 repeat: matches 1603..2053 of consensus"
 10921..11230
 /note="AluSg repeat: matches 1..310 of consensus"
 11231..14323
 /note="L1MB7 repeat: matches 2053..4281 of consensus"
 14324..14604
 /note="L1MC2 repeat: matches 6052..6330 of consensus"

misc_feature

repeat_region

misc_feature

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               19411..20261
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               /evidence=not_experimental
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               21866..22366
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               24137..24358
repeat_region /note="MIR repeat: matches 6..241 of consensus"
               24511..24618
repeat_region /note="MIR repeat: matches 119..229 of consensus"
               25107..27565
repeat_region /note="L1PA7 repeat: matches 3679..6145 of consensus"
               29303..29618
repeat_region /note="MER1B repeat: matches 2..337 of consensus"
               30113..30371
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seq_name: gb_htg4:AC011241

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LOCUS      AC011241 187493 bp DNA HTG 29-OCT-1999
DEFINITION Homo sapiens chromosome 2 clone NH0467P13 map unknown, WORKING
DRAFT SEQUENCE, in unordered pieces.
ACCESSION  AC011241
VERSION     AC011241.2 GI:6139206
KEYWORDS    HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE      human.
ORGANISM    Homo sapiens
             Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
             Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE   1 (bases 1 to 187493)
AUTHORS     Sulston,J.E. and Waterston,R.
TITLE       Toward a complete human genome sequence
JOURNAL     Genome Res. 8 (11), 1097-1108 (1998)
MEDLINE     99063792
REFERENCE   2 (bases 1 to 187493)
AUTHORS     Waterston,R.
TITLE       The sequence of Homo sapiens unknown clone NH0467P13
JOURNAL     Unpublished
REFERENCE   3 (bases 1 to 187493)
AUTHORS     Waterston,R.H.
TITLE       Direct Submission
JOURNAL     Submitted (04-OCT-1999) Genome Sequencing Center, Washington
             University School of Medicine, 4444 Forest Park Parkway, St. Louis,
             MO 63108, USA
             On Oct 29, 1999 this sequence version replaced gi:6007883.
COMMENT     SUBMITTED BY: WUGSC
             Genome Sequencing Center
             Department of Genetics
             Washington University
             St. Louis MO 63108, USA
             http://genome.wustl.edu/gsc
             mailto:sapiens@watson.wustl.edu

NOTICE: This 'working draft' quality sequence may consist of
several contigs from automated sequence assembly concatenated
together. No attempt has been made to order or orient the contigs
relative to one another correctly before concatenating. At each
location in the sequence where contigs have been joined, several
consecutive Ns may have been inserted.

The attached annotation was produced using a purely automated
procedure.
* NOTE: This is a 'working draft' sequence.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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                     624..932
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seq_name: gb_ba2:AF026541
seq_documentation_block:
LOCUS AF026541 1692 bp DNA BCT 30-OCT-1998
DEFINITION Mycobacterium tuberculosis CeoB (ceoB) gene, complete cds; and CeoC (ceoC) gene, partial cds.
ACCESSION AF026541
VERSION AF026541.1 GI:2582553
SOURCE Mycobacterium tuberculosis.
ORGANISM Mycobacterium tuberculosis
Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
Actinomycetales; Corynebacterineae; Mycobacteriaceae;
Mycobacterium; Mycobacterium tuberculosis complex.
1 (bases 1 to 1692)
Chen, P. and Bishai, W. R.
Novel selection for isoniazid (INH) resistance genes supports a
role for NAD+-binding proteins in mycobacterial INH resistance
Infect. Immun. 66 (11), 5099-5106 (1998)
99003115
2 (bases 1 to 1692)
Chen, P. and Bishai, W. R.
Direct Submission
Submitted (23-SEP-1997) Molecular Microbiology and Immunology,
Johns Hopkins University, 615 N. Wolfe Street, Baltimore, MD 21205,
USA
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phenotype of oxyR deletion mutant of E. coli"
/codon_start=1
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/db_xref="GI:2582554"
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BASE COUNT 337 a 544 c 550 g 261 t
ORIGIN

alignment_scores:
Quality: 36.00 Length: 12
Ratio: 4.000 Gaps: 0
Percent Similarity: 75.000 Percent Identity: 66.667

OM of: US-08-653-294-19 to: N_Geneseq_36:* out_format : pfs
 Date: Feb 8, 2000 1:28 PM
 About: Results were produced by the GenCore software, version 4.5,
 Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:
 -MODEL=framed-p2n.model -DEV=xlpx
 -Q/cgml_1/USPTO.spool/US08653294/runat_04022000_160701_15807/app_query.fasta.1
 -DB=N_Geneseq_36 -OFMT=fastap -SUFFIX=ring -GAPOP=12.000
 -GAPEXT=4.000 -MINMATCH=0.100 -LOOPEXT=0.000 -LOOPEXT=0.000
 -GAPOP=4.500 -GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500
 -XGAPOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=blomsum62
 -TRANS=human40.cdi -LIST=45 -DOALIGN=200 -THR_SCORE=pct
 -ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -MINLEN=0
 -MAXLEN=1000000 -USER=US08653294 -NCPU=6 -ICPU=3 -NO_XLPXY -WAIT
 -THREADS=1

Search information block:

Query: US-08-653-294-19
 Query length: 12
 Database: N_Geneseq_36:*
 Database sequences: 3111585
 Database length: 125096042
 Search time (sec): 590.520000

score_list:

Sequence	Strd Orig	ZScore	EScore	Len	Documentation
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N_Geneseq_36:T76573	36.00	116.81	66.13	1728	Pig alpha 1-6 fucosyltransferase
N_Geneseq_36:T1310	35.00	142.22	2.54	58	D-A42d RNA sequence. L-nucleic
N_Geneseq_36:T74678	35.00	112.44	115.91	1847	Staphylococcus aureus contig 8
N_Geneseq_36:T66241	35.00	111.84	125.23	1981	Romaine lettuce violaxanthin d
N_Geneseq_36:T644280	34.00	113.16	105.69	1092	Pseudomonas cepacia DSM 3401
N_Geneseq_36:T12978	34.00	108.13	201.46	1959	Enterococcus faecalis genome c
N_Geneseq_36:T13340	34.00	106.14	260.08	2469	Enterococcus faecalis genome c
N_Geneseq_36:T19375	33.00	100.65	525.78	3003	Hereditary multiple exostose a
N_Geneseq_36:T13357	33.00	97.94	744.22	4114	Enterococcus faecalis genome c
N_Geneseq_36:T13127	33.00	89.16	2.3e+03	11410	Enterococcus faecalis genome c
N_Geneseq_36:T00049	32.00	115.67	76.58	337	Hepatitis GB virus (HGBV) clone
N_Geneseq_36:T00127	32.00	115.67	76.58	337	Hepatitis GB virus (HGBV) clone
N_Geneseq_36:T41700	32.00	103.75	353.42	1347	Brugia pahangi beta-tubulin c
N_Geneseq_36:T12225	32.00	103.59	360.38	1371	Octopus rhodopsin membrane p
N_Geneseq_36:T21309_11	32.00	65.86	4.4e+04	110000	Continuation (12 of 17) of
N_Geneseq_36:T80367	31.00	126.72	18.56	60	Heteropolymer oligo used in immu
N_Geneseq_36:T80372	31.00	126.30	19.59	63	Phosphorylated oligo used in a h
N_Geneseq_36:T27693	31.00	118.01	56.70	165	PGEM32f target fragment. Cleava
N_Geneseq_36:T27694	31.00	116.10	72.44	206	PGEM32f target fragment. Cleava
N_Geneseq_36:T70340	31.00	116.10	72.44	206	DNA substrate for 5' nuclease.
N_Geneseq_36:T76640	31.00	116.10	72.44	206	Duplex sequence used as 5' nucl
N_Geneseq_36:T53874	31.00	116.10	72.44	206	Nucleotide sequence of the 5' n
N_Geneseq_36:T65800	31.00	116.10	72.44	206	5' Nuclease substrate DNA. Det
N_Geneseq_36:T63426	31.00	116.10	72.44	206	206-mer duplex used as a substr
N_Geneseq_36:T080775	31.00	115.23	81.02	228	5' Nuclease substrate. 5' Nucle
N_Geneseq_36:T16481	31.00	110.18	154.85	410	DNA encoding a Bacillus thurin
N_Geneseq_36:T16485	31.00	110.18	154.85	410	DNA encoding a Bacillus thurin
N_Geneseq_36:T21079	31.00	109.71	164.47	433	Polynucleotide sequence from th
N_Geneseq_36:T28565	31.00	103.80	351.23	861	Bacterial antibiotic resistance
N_Geneseq_36:T24873	31.00	101.07	498.30	1182	Alpha-ald gene. Accelerated b
N_Geneseq_36:T20675	31.00	100.75	518.81	1226	Polynucleotide sequence from t
N_Geneseq_36:T82205	31.00	100.50	536.12	1263	Nad B gene encoding L aspartat
N_Geneseq_36:T80877	31.00	99.61	600.66	1400	Sequence encoding Serratia ph
N_Geneseq_36:T70130	31.00	99.39	618.21	1437	Max-interacting protein coding
N_Geneseq_36:T03064	31.00	96.58	885.54	1990	Encodes Babesia bovis 60kD im
N_Geneseq_36:T18995	31.00	96.58	885.54	1990	Babesia merosioite surface prot
N_Geneseq_36:T39653	31.00	93.75	1.3e+03	2765	Renal cancer associated gene.
N_Geneseq_36:T12366	31.00	93.06	1.4e+03	2996	Gene encoding enzyme with stan
N_Geneseq_36:T1822	31.00	92.23	1.5e+03	3300	Mutant Aspergillus oryzae DEB
N_Geneseq_36:T59579	31.00	92.19	1.6e+03	3315	Penicillium chrysogenum acetam

N_Geneseq_36:T16517 + 31.00 91.84 1.6e+03 3453 ! DNA encoding a Bacillus thu
 N_Geneseq_36:T61556 + 31.00 91.31 1.7e+03 3672 ! Human signal mediator prote
 N_Geneseq_36:T38466 + 31.00 90.87 1.8e+03 3867 ! DNA encoding a Bacillus thu
 N_Geneseq_36:T20635 - 31.00 89.67 2.1e+03 4444 ! Polynucleotide sequence fro

seq_name: N_Geneseq_36:T26331

seq_documentation_block:

ID T26331 standard; cDNA to mRNA; 382 BP.
 AC T26331;
 DT 16-OCT-1996 (first entry)
 DE Human gene signature HUMG08571.
 KW Gene signature; messenger RNA; mRNA; relative abundance; frequency;
 KW human; cloning; mapping; non-biased library; diagnosis; detection;
 KW cell typing; abnormal cell function; ss.
 OS Homo sapiens.
 PN W09514772-A1.
 PD 01-JUN-1995.
 PF 11-NOV-1994; J01916.
 PR 12-NOV-1993; JP-353504.
 PA (MATS/) MATSUBARA K.
 PI (OKUB/) OKUBO K.
 PI Matsubara K. Okubo K;
 DR WPI; 95-206931/27.
 PT Identifying gene signatures in 3'-directed human cDNA library - e.g.
 PT for diagnosis of abnormal cell function, by preparing cDNA that
 PT reflects relative abundance of corresp. mRNA in specific human
 PT tissues
 PS Claim 1: Page 2058-2059; 2245pp; Japanese.
 CC A single-stranded DNA (or its complementary strand or the corresp.
 CC double-stranded DNA) which comprises one of the 7837 "GS" sequences
 CC given in T19001-T26837 and which is able to hybridise to part of
 CC human genomic DNA, cDNA or mRNA is claimed. The GS (Gene Signature)
 CC sequences were obtained from 3'-directed cDNA libraries prepared
 CC from various human tissues; synthesis of cDNA was initiated from the
 CC 3'-end of mRNA by using poly(rI) as the sole primer. Since the 3'-
 CC untranslated sequence is unique to a particular mRNA species, almost
 CC all the 3'-oriented cDNAs hybridise with specific mRNAs. Each library
 CC is constructed so as to reflect accurately the relative abundance of
 CC different mRNAs in the particular tissue from which it was derived.
 CC The appearance frequency of a given GS in a cDNA library can be
 CC determined (esp. using primers and probes derived from the GS
 CC sequences) as a means of diagnosing abnormal cell function or for
 CC recognising different cell types.
 SQ Sequence 382 BP; 118 A; 72 C; 85 G; 102 T;

alignment_scores:

Quality: 36.00 Length: 12
 Ratio: 4.000 Gaps: 0
 Percent Similarity: 75.000 Percent Identity: 58.333

alignment_block:

US-08-653-294-19 x T26331 ..

Align seg 1/1 to: T26331 from: 1 to: 382

1 TvrArgLeuAlaIle***ArgileAlaLeuArgTyr 12
 ||||| :||:|||||||
 36 TACAGCGGTGACTACCGCGGCTGCGCTCAGATAT 71

seq_name: N_Geneseq_36:T76573

seq_documentation_block:

ID T76573 standard; cDNA to mRNA; 1728 BP.
 AC T76573;
 DT 05-MAR-1998 (first entry)
 DE Pig alpha 1-6 fucosyltransferase gene.
 KW Alpha 1-6 fucosyltransferase; enzyme; pig; human; fucose transfer;
 KW guanosine diphosphate; sugar chain synthesis; modification; antibody;
 KW GlcNAc; cancer diagnosis; ss.
 OS Sus scrofa.
 FH Key Location/Qualifiers

```

FT CDS 1. .1728
FT /tag= a
FT WO9727303-A1.
FT 31-JUL-1997;
FT 23-JAN-1997; J00171.
FT 22-JUL-1996; JP-192260.
FT 24-JAN-1996; JP-010365.
FT 21-JUN-1996; JP-161648.
FT 24-JUN-1996; JP-162813.
FT (TOYM ) TOYO BOSEKI KK.
FT Shiba T., Taniguchi N, Uozumi N, Yanagidani S;
FT WPI: 97-393690/36.
FT P-PSDB; W2124.
FT Human or pig alpha 1-6 fucosyl:transferase and DNA encoding it - for
FT synthesis and modification of sugar chains and used as an antigen
FT for production of diagnostic antibodies
FT Claim 5; Page 30-34; 61pp; Japanese.
FT T76573 and T76574 represent the coding sequences for the pig and human
FT alpha 1-6 fucosyltransferases of the invention, respectively. The encoded
FT enzyme transfers fucose from guanosine diphosphate to the 6-hydroxyl
FT group of the GlcNAc nearest to R in the receptor molecule: (GlcNAc
FT 1-2Manalpa 1-6)(GlcNAc 1-2Manalpa 1-3)Manbeta 1-4GlcNAc 1-2Manalpa
FT 1-4GlcNAc-R to give (GlcNAc 1-2Manalpa 1-6)(GlcNAc 1-2Manalpa
FT 1-3)Manbeta 1-4GlcNAc 1-4(Fucalpa 1-6)GlcNAc-R. It has an optimum pH
FT of about 7.0 (pig) or 7.5 (human), and is stable over the pH range 4-10
FT after 5 hours at 4 degrees C. The optimum working temperature of the
FT enzyme is 30-37 degrees C. A bivalent metal is not required for activity
FT of the enzyme, and the enzyme is not inhibited in the presence of 5 mM
FT EDTA. The enzyme is useful in the synthesis and modification of sugar
FT chains, and as antigen for the production of antibodies recognising the
FT enzyme. The antibodies can be used for the diagnosis of cancer and other
FT diseases.
FT Sequence 1728 BP; 521 A; 362 C; 419 G; 426 T;

alignment_scores:
Quality: 36.00 Length: 11
Ratio: 3.273 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 72.727

alignment_block:
US-08-653-294-19 x T76573 ..
Align seg 1/1 to: T76573 from: 1 to: 1728

2 ArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||
723 CGCCTTGGAATCTCACAATGGCGCTACGCTAC 755

seq_name: N_Geneseq_36.V11310

seq_documentation_block:
ID V11310 standard; RNA; 58 BP.
AC V11310;
DT 31-JUL-1998 (first entry)
DE D-A42d RNA sequence.
KW L-nucleic acid; target; D-nucleic acid; optical antipode; therapy;
KW diagnosis; biosensor; herbicide; food additive; analysis; perfume;
KW flavouring; cosmetic; purification; dialysis; separation; tumour;
KW viral infection; bacterial infection; ss.
OS Unknown.
FH Key Location/Qualifiers
FT misc_binding 1. .8
FT /tag= a
FT /note= "Binds to nucleotides 51. .58"
FT
FT misc_binding 9. .10
FT /tag= b
FT /note= "Binds to nucleotides 46. .47"
FT
FT stem_loop 16. .26
FT /tag= c
FT stem_loop 29. .45
FT /tag= d
FT misc_binding 46. .47

FT /tag= e
FT /note= "Binds to nucleotides 9. .10"
FT
FT misc_binding 51. .58
FT /tag= f
FT /note= "Binds to nucleotides 1. .8"
FT
WO9808856-A2.
05-MAR-1998.
29-AUG-1997; EP04726.
30-AUG-1996; EP-113953.
(BALD/) BALD R.
(ERDM/) ERDMANN V A.
(FUERK/) FUERSTE J P.
Bald R, Erdmann VA, Fuerste JP;
WPI: 98-179376/16.
L-nucleic acids that bind target molecules selectively - isolated by
screening D-nucleic acids against optical antipode, useful in
therapy and diagnosis
Disclosure: Fig 13; 89pp; German.
This sequence is used in a method which results in the production of
L-nucleic acids able to bind to a target molecule. The method involves
generating a heterogeneous population of D-nucleic acids then treating
them with optical antipodes. Such nucleic acids are useful in therapy
and diagnosis, as e.g. biosensors, herbicides, food additives, in the
analysis of perfumes or flavourings and in cosmetics, for formulating
e.g. sunscreens or anti-wrinkle creams. They may stimulate or inhibit
the function of the target molecule and can be coupled to markers or
cytotoxins, carriers for affinity purification of the target molecule,
including separation of enantiomers or elimination of enantiomeric
impurities, or they can be used for purification of cellular factors or
cells, e.g. the separation of toxic components by dialysis. The nucleic
acids can be used in therapeutic applications for e.g. for tumours,
viral and bacterial infections and for hypertension. The method allows
selection and evaluation of high affinity nucleic acids. D-nucleic acids
and optical antipodes of the target are more stable under physiological
conditions than compounds with the natural configuration (particularly
in the case of RNA). This stability means that potentially harmful
metabolites are not formed and immune responses are not induced.
Sequence 58 BP; 13 A; 14 C; 13 G; 18 U;

alignment_scores:
Quality: 35.00 Length: 12
Ratio: 3.889 Gaps: 0
Percent Similarity: 75.000 Percent Identity: 50.000

alignment_block:
US-08-653-294-19 x V11310 ..
Align seg 1/1 to: V11310 from: 1 to: 58

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||
6 UACCGCAAAAGCGUUUUGCGCAUACCAUUGCUU 41

seq_name: N_Geneseq_36.V74678

seq_documentation_block:
ID V74678 standard; DNA; 1847 BP.
AC V74678;
DT 16-MAR-1999 (first entry)
DE Staphylococcus aureus contig SEQ ID #367.
KW Computer readable medium; vaccine; S.aureus infection; immunodetection;
KW cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;
KW skin infection; surgical wound infection; scalded skin syndrome;
KW toxic shock syndrome; ds.
OS Staphylococcus aureus.
FH Key Location/Qualifiers
FT misc_feature 1381. .1440
FT /tag= a
FT /note= "these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"

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PN EP-786519-A2.
 PD 30-JUL-1997.
 PF 07-JAN-1997; 100117.
 PR 05-JAN-1996; US-009861.
 PA (HUMA-) HUMAN GENOME SCI INC.
 PI Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA,
 PI Rosen CA;
 DR WPI: 97-374922/35.
 PT Polynucleotide(s) and proteins derived from *Staphylococcus aureus* -
 PT stored on computer readable medium and used in the production of
 PT anti-S aureus vaccines
 PS Claim 1: Page 1260-1261; 3271pp; English.
 CC This sequence represents one of 5191 *Staphylococcus aureus* DNA sequences
 CC of the invention. The DNA sequences are recorded on a computer readable
 CC medium, preferably selected from a floppy or hard disk, random access
 CC memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
 CC the S aureus DNA sequences allows putative functions to be assigned so
 CC that protein-encoding or regulatory regions of commercial, therapeutic or
 CC industrial importance can be obtained. Specifically, sequences which are
 CC likely to encode antigens have been identified and these polypeptides can
 CC be used in a vaccine composition against S aureus infection. The
 CC polypeptides can also be used in a kit for the immunodetection of
 CC S aureus in a sample. S aureus is implicated in numerous human diseases,
 CC including cellulitis, eyelid infections, food poisoning, osteomyelitis,
 CC skin and surgical wound infections, scalded skin syndrome, toxic shock
 CC syndrome, etc. Organisms transformed with the DNA sequences can be used
 CC for recombinant production of the polypeptides. The new DNA sequences
 CC (and their fragments) are useful as primers or probes for isolating
 CC homologues of any of the S aureus DNA sequences contained on the
 CC computer readable medium.
 SQ Sequence 1847 BP; 627 A; 257 C; 334 G; 567 T;

alignment_scores:
 Quality: 35.00 Length: 10
 Ratio: 3.500 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 80.000

alignment_block:
 US-08-653-294-19 x V74678 ..

Align seg 1/1 to: V74678 from: 1 to: 1847

3 LeuAlaIle***ArgIleAlaLeuArgTyr 12
 :::::::::::::::::::::::::::::::
 101 GTTGCTATTCTAGATTGCATTGGCTTAT 130

seq_name: N_Geneseq_36:T66241

seq_documentation_block:
 ID T66241 standard; cdna; 1981 BP.
 AC T66241;
 DT 28-JUL-1997 (first entry)
 DE Romaine lettuce violaxanthin de-epoxidase cDNA.
 KW Violaxanthin de-epoxidase; VDE; light; photosensitivity;
 KW photoprotection; transgenic plant; zeaxanthin; antheraxanthin;
 KW xanthophyll; lettuce; ss.
 OS Lactuca sativa L. cv. romaine.
 FH Key Location/Qualifiers
 FT misc_difference 26..29
 FT /tag= a
 FT /note= "bases 26-29 are illegible in Fig 1"
 FT misc_difference 66..72
 FT /tag= b
 FT /note= "bases 66-72 are illegible in Fig 1"
 FT misc_difference 105..110
 FT /tag= c
 FT /note= "bases 105-110 are illegible in Fig 1"
 FT misc_difference 147..149
 FT /tag= d
 FT /note= "bases 147-149 are illegible in Fig 1"
 FT misc_difference 186..189
 FT /tag= e

FT misc_difference 226..227
 FT /tag= f
 FT /note= "bases 226-227 are illegible in Fig 1"
 FT cds 235..1656
 FT /tag= g
 FT transit_peptide 235..609
 FT /tag= h
 FT mat_peptide 610..1653
 FT /tag= i
 PN W09717447-A2.
 PD 15-MAY-1997.
 PF 07-NOV-1996; U18291.
 PR 07-NOV-1995; US-006315.
 PR 06-AUG-1996; US-023502.
 PA (CALJ) CALGENE INC.
 PI Bugos RC, Rockholm DC, Yamamoto HY;
 DR WPI: 97-281036/25.
 PT P-PSDB; W09874.
 CC A CDNA clone (T66241) codes for the 55 kDa violaxanthin de-epoxidase
 CC (VDE) (W09874) of romaine lettuce. VDE was purified from romaine
 CC lettuce chloroplasts and 2 tryptic peptides were used to develop
 CC primers (see also T66244-45), which amplified a partial VDE
 CC sequence. The amplified sequence was then used to screen a lettuce
 CC cDNA library, and the 1981 bp DNA sequence was identified. VDE
 CC nucleic acids (see also T66242-43), in sense or antisense
 CC orientation, can be used in genetic constructs, pref. also contg. a
 CC plastid translocation sequence, to modify VDE levels in plants.
 CC Increased levels result in the plant being tolerant of increased
 CC light and therefore more productive and/or more resistant to
 CC disease. Underexpression of VDE increases photosynthetic
 CC efficiency under low light. The photosensitivity of a range of
 CC crops, trees and ornamentals can be modified.
 SQ Sequence 1981 BP; 608 A; 337 C; 433 G; 577 T;

alignment_scores:
 Quality: 35.00 Length: 10
 Ratio: 3.889 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 90.000

alignment_block:
 US-08-653-294-19 x T66241 ..

Align seg 1/1 to: T66241 from: 1 to: 1981

3 LeuAlaIle***ArgIleAlaLeuArgTyr 12
 :::::::::::::::::::::::::::::::
 434 CTAGCCATTGCAAGGATAAATCTCAGATAT 463

seq_name: N_Geneseq_36:Q44280

seq_documentation_block:
 ID Q44280 standard; DNA; 1092 BP.
 AC Q44280;
 DT 09-DEC-1993 (first entry)
 DE Pseudomonas cepacia DSM 3401 lipD gene.
 KW Lipase; LipD; lipase modulator; lmbD; chaperone molecule;
 KW lipolysis; detergent; ss.
 OS Pseudomonas cepacia.
 FH Key Location/Qualifiers
 FT misc_difference 391
 FT /tag= a
 FT misc_difference 480
 FT /tag= b
 FT misc_difference 481
 FT /tag= c
 FT misc_difference 712..714
 FT /tag= d
 FT misc_difference 734
 FT /tag= e

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FT WO9313200-A.
PN 08-JUL-1993.
PF 18-DEC-1992; DK0391.
PR 20-DEC-1991; WO-DK0402.
PA (NOVO ) NOVO-NORDISK AS.
PI Buckley CM, Diderichsen BK, Hobson A, Joergensen ST;
PI McConnell DJ;
DR WPI: 93-227318/28.
DR P-PSDB; R39396.
PT Prep. of active lipase in high quantities - by subjecting to
PT denaturation and restructuring in presence of chaperone molecule
PS Example 10; Page 43; 78pp; English.
CC Two genes were cloned and sequenced from Pseudomonas cepacia DSM
CC 3401. The genes were designated lipD (Q44280) and lipB (Q44281) and
CC they code for a lipase and a lipase modulator protein, respectively.
CC Due to the extreme GC content of the DNA, the sequence was difficult
CC to determine (hence the "Others" in the sequence). The lipD start
CC codon is positioned 3 bp downstream of the lipD stop codon. LipD and
CC LipB were found to be homologous to LipA and Lima, respectively. In
CC denaturation/renaturation experiments, Lima chaperone protein was
CC able to produce active LipD.
SQ Sequence 1092 BP; 188 A; 353 C; 378 G; 166 T;

alignment_scores:
  Quality: 34.00 Length: 10
  Ratio: 4.250 Gaps: 0
Percent Similarity: 80.000 Percent Identity: 70.000

alignment_block:
US-08-653-294-19 x Q44280/rev ..
Align seg 1/1 to reverse of: Q44280 from: 1 to: 1092

3 LeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||: ||| |||||
960 CTGGCCGTACAGCGCTGCACTTCGATAC 931

seq_name: N_Geneseq_36:X12978
seq_documentation_block:
ID X12978 standard; DNA; 1959 BP.
AC X12978;
DT 19-MAR-1999 (first entry)
DE Enterococcus faecalis genome contig SEQ ID NO:41.
KW Enterococcus faecalis; contig; detection; Enterococcal infection;
KW vaccine; attenuation; computer readable medium; ds.
OS Enterococcus faecalis.
PN WO9850555-A2.
PD 12-NOV-1998.
PF 04-MAY-1998; U08985.
PR 14-NOV-1997; US-066009.
PR 06-MAY-1997; US-044031.
PR 16-MAY-1997; US-046655.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Dillon PJ, Kunsch CA;
DR WPI: 99-045171/04.
PT New isolated Enterococcus faecalis polynucleotides and polypeptides
PT - used to develop products for the detection of Enterococcus and for
PT use in vaccines for prevention or attenuation of Enterococcus
PT infection.
PS Claim 1; Page 413-414; 2084pp; English.
CC A computer readable medium has been developed which has recorded on it
CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.
CC X12938 to X13919 represent these nucleotide sequences which are primary
CC nucleotide sequences, also known as contigs. The computer-based system
CC can identify fragments of the Enterococcus faecalis genome with
CC commercial importance. The products can be used to detect the presence
CC of Enterococcus faecalis in samples. They can also be used for
CC diagnosing Enterococcal infection in an animal and monitoring
CC progression of disease, and for identifying agents which can be used to
CC modulate the growth or pathogenicity of Enterococcus faecalis, or
CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.
SQ Sequence 2469 BP; 771 A; 471 C; 483 G; 738 T;

alignment_scores:
  Quality: 34.00 Length: 12
  Ratio: 3.091 Gaps: 0
Percent Similarity: 91.667 Percent Identity: 50.000

alignment_block:
US-08-653-294-19 x X13540/rev ..
Align seg 1/1 to reverse of: X13540 from: 1 to: 2469

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||: ||| |||||
1358 TATATACCTGCTATATTAGATTTTCTTCGGTAT 1393

seq_name: N_Geneseq_36:X13540
seq_documentation_block:
ID X13540 standard; DNA; 2469 BP.
AC X13540;
DT 19-MAR-1999 (first entry)
DE Enterococcus faecalis genome contig SEQ ID NO:603.
KW Enterococcus faecalis; contig; detection; Enterococcal infection;
KW vaccine; attenuation; computer readable medium; ds.
OS Enterococcus faecalis.
PN WO9850555-A2.
PD 12-NOV-1998.
PF 04-MAY-1998; U08985.
PR 14-NOV-1997; US-066009.
PR 06-MAY-1997; US-044031.
PR 16-MAY-1997; US-046655.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Dillon PJ, Kunsch CA;
DR WPI: 99-045171/04.
PT New isolated Enterococcus faecalis polynucleotides and polypeptides
PT - used to develop products for the detection of Enterococcus and for
PT use in vaccines for prevention or attenuation of Enterococcus
PT infection.
PS Claim 1; Page 1853-1855; 2084pp; English.
CC A computer readable medium has been developed which has recorded on it
CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.
CC X12938 to X13919 represent these nucleotide sequences which are primary
CC nucleotide sequences, also known as contigs. The computer-based system
CC can identify fragments of the Enterococcus faecalis genome with
CC commercial importance. The products can be used to detect the presence
CC of Enterococcus faecalis in samples. They can also be used for
CC diagnosing Enterococcal infection in an animal and monitoring
CC progression of disease, and for identifying agents which can be used to
CC modulate the growth or pathogenicity of Enterococcus faecalis, or
CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.
SQ Sequence 2469 BP; 771 A; 471 C; 483 G; 738 T;

alignment_scores:
  Quality: 34.00 Length: 12
  Ratio: 3.091 Gaps: 0
Percent Similarity: 91.667 Percent Identity: 50.000

alignment_block:
US-08-653-294-19 x X13540/rev ..
Align seg 1/1 to reverse of: X13540 from: 1 to: 2469

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||: ||| |||||
1358 TATATACCTGCTATATTAGATTTTCTTCGGTAT 1393

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CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.
SQ Sequence 1959 BP; 542 A; 341 C; 322 G; 749 T;

alignment_scores:
  Quality: 34.00 Length: 12
  Ratio: 3.778 Gaps: 0
Percent Similarity: 75.000 Percent Identity: 66.667

alignment_block:
US-08-653-294-19 x X12978 ..
Align seg 1/1 to: X12978 from: 1 to: 1959

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||: ||| |||||
1358 TATATACCTGCTATATTAGATTTTCTTCGGTAT 1393

seq_name: N_Geneseq_36:X13540
seq_documentation_block:
ID X13540 standard; DNA; 2469 BP.
AC X13540;
DT 19-MAR-1999 (first entry)
DE Enterococcus faecalis genome contig SEQ ID NO:603.
KW Enterococcus faecalis; contig; detection; Enterococcal infection;
KW vaccine; attenuation; computer readable medium; ds.
OS Enterococcus faecalis.
PN WO9850555-A2.
PD 12-NOV-1998.
PF 04-MAY-1998; U08985.
PR 14-NOV-1997; US-066009.
PR 06-MAY-1997; US-044031.
PR 16-MAY-1997; US-046655.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Dillon PJ, Kunsch CA;
DR WPI: 99-045171/04.
PT New isolated Enterococcus faecalis polynucleotides and polypeptides
PT - used to develop products for the detection of Enterococcus and for
PT use in vaccines for prevention or attenuation of Enterococcus
PT infection.
PS Claim 1; Page 1853-1855; 2084pp; English.
CC A computer readable medium has been developed which has recorded on it
CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.
CC X12938 to X13919 represent these nucleotide sequences which are primary
CC nucleotide sequences, also known as contigs. The computer-based system
CC can identify fragments of the Enterococcus faecalis genome with
CC commercial importance. The products can be used to detect the presence
CC of Enterococcus faecalis in samples. They can also be used for
CC diagnosing Enterococcal infection in an animal and monitoring
CC progression of disease, and for identifying agents which can be used to
CC modulate the growth or pathogenicity of Enterococcus faecalis, or
CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.
SQ Sequence 2469 BP; 771 A; 471 C; 483 G; 738 T;

alignment_scores:
  Quality: 34.00 Length: 12
  Ratio: 3.091 Gaps: 0
Percent Similarity: 91.667 Percent Identity: 50.000

alignment_block:
US-08-653-294-19 x X13540/rev ..
Align seg 1/1 to reverse of: X13540 from: 1 to: 2469

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||: ||| |||||
1358 TATATACCTGCTATATTAGATTTTCTTCGGTAT 1393

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1322 TTCAGTCTTCTTAACGGCGGTTTCGCTCGGTAT 1287
seq_name: N_Geneseq_36:V19375
seq_documentation_block:
ID V19375 standard; CDNA: 3003 BP.
AC V19375;
DT 20-AUG-1998 (first entry)
DE Hereditary multiple exostoses associated EXT2 gene isoform encoding cDNA.
KW Hereditary multiple exostoses; EXT2; chondrosarcoma; human; isoform;
KW treatment; ss.
OS Homo sapiens.
FH Key Location/Qualifiers
FT CDS 167..2353
FT /tag= a
FT /product= "EXT2 gene isoform"
PN EP-837127-A2.
PD 22-APR-1998.
PF 26-AUG-1997; 306503.
PR 21-OCT-1996; CN-121928.
PA (UVHU-) UNIV HUNAN MEDICAL.
PI Deng HX, Fan CH, Xia J;
DR P-PSDB; W44851.
DR WPI: 98-219110/20.
PT Cloned human EXT2 gene - associated with hereditary multiple
PT exostoses or chondrosarcoma
PS Claim 4: Pages 28-31; 31pp; English.
CC This cDNA encodes an isoform of the EXT2 gene associated with hereditary
CC multiple exostoses and chondrosarcoma. The polynucleotide is an isoform
CC of the EXT2 gene described in Nature Genet., 14, 25, 1996. The
CC polynucleotide can be used in the detection and treatment of EXT2-related
CC diseases, and to identify compounds which activate or inhibit receptors
CC for the encoded polypeptide. The polypeptide can be recombinantly
CC produced by transforming or transfecting a cell with a vector containing
CC the encoding nucleic acid.
SQ Sequence 3003 BP; 751 A; 730 C; 766 G; 756 T;

alignment_scores:
Quality: 33.00 Length: 11
Ratio: 3.000 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 54.545

alignment_block:
US-08-653-294-19 x V19375 ..
Align seg 1/1 to: V19375 from: 1 to: 3003
1 TyArgLeuAlaIle***ArgIleAlaLeuArg 11
||||:||||:||||:||||:||||:||||:
1170 TACAAGCTGCTGTCGCCGGTGTGTCATTGCAG 1202

seq_name: N_Geneseq_36:X13357
seq_documentation_block:
ID X13357 standard; DNA: 4114 BP.
AC X13357;
DT 19-MAR-1999 (first entry)
DE Enterococcus faecalis genome contig SEQ ID NO:420.
KW Enterococcus faecalis; contig; detection; Enterococcal infection;
KW vaccine; attenuation; computer readable medium; ds.
OS Enterococcus faecalis.
PN WO9850555-A2.
PD 12-NOV-1998.
PF 04-MAY-1998; U08985.
PR 14-NOV-1997; US-066009.
PR 06-MAY-1997; US-044031.
PR 16-MAY-1997; US-046655.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Dillon PJ, Kunsch CA;
DR WPI: 99-045171/04.
PT New isolated Enterococcus faecalis polynucleotides and polypeptides
PT - used to develop products for the detection of Enterococcus and for
PT use in vaccines for prevention or attenuation of Enterococcus
PT infection.
PS Claim 1: Page 1021-1027; 2084pp; English.
CC A computer readable medium has been developed which has recorded on it
CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.
CC X12938 to X13919 represent these nucleotide sequences which are primary
CC nucleotide sequences, also known as contigs. The computer-based system
CC can identify fragments of the Enterococcus faecalis genome with
CC commercial importance. The products can be used to detect the presence
CC of Enterococcus faecalis in samples. They can also be used for
CC diagnosing Enterococcal infection in an animal and monitoring
CC progression of disease, and for identifying agents which can be used to
CC modulate the growth or pathogenicity of Enterococcus faecalis, or
CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.

PT - used to develop products for the detection of Enterococcus and for
PT use in vaccines for prevention or attenuation of Enterococcus
PT infection.
PS Claim 1: Page 1639-1641; 2084pp; English.
CC A computer readable medium has been developed which has recorded on it
CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.
CC X12938 to X13919 represent these nucleotide sequences which are primary
CC nucleotide sequences, also known as contigs. The computer-based system
CC can identify fragments of the Enterococcus faecalis genome with
CC commercial importance. The products can be used to detect the presence
CC of Enterococcus faecalis in samples. They can also be used for
CC diagnosing Enterococcal infection in an animal and monitoring
CC progression of disease, and for identifying agents which can be used to
CC modulate the growth or pathogenicity of Enterococcus faecalis, or
CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.
SQ Sequence 4114 BP; 1334 A; 661 C; 889 G; 1226 T;

alignment_scores:
Quality: 33.00 Length: 11
Ratio: 3.667 Gaps: 0
Percent Similarity: 81.818 Percent Identity: 54.545

alignment_block:
US-08-653-294-19 x X13357 ..
Align seg 1/1 to: X13357 from: 1 to: 4114
2 ArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
||||:||||:||||:||||:||||:
1110 AGAATGCTTACGACGATTAGCTAGTCGTAT 1142

seq_name: N_Geneseq_36:X13127
seq_documentation_block:
ID X13127 standard; DNA: 11410 BP.
AC X13127;
DT 19-MAR-1999 (first entry)
DE Enterococcus faecalis genome contig SEQ ID NO:190.
KW Enterococcus faecalis; contig; detection; Enterococcal infection;
KW vaccine; attenuation; computer readable medium; ds.
OS Enterococcus faecalis.
PN WO9850555-A2.
PD 12-NOV-1998.
PF 04-MAY-1998; U08985.
PR 14-NOV-1997; US-066009.
PR 06-MAY-1997; US-044031.
PR 16-MAY-1997; US-046655.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Dillon PJ, Kunsch CA;
DR WPI: 99-045171/04.
PT New isolated Enterococcus faecalis polynucleotides and polypeptides
PT - used to develop products for the detection of Enterococcus and for
PT use in vaccines for prevention or attenuation of Enterococcus
PT infection.
PS Claim 1: Page 1021-1027; 2084pp; English.
CC A computer readable medium has been developed which has recorded on it
CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.
CC X12938 to X13919 represent these nucleotide sequences which are primary
CC nucleotide sequences, also known as contigs. The computer-based system
CC can identify fragments of the Enterococcus faecalis genome with
CC commercial importance. The products can be used to detect the presence
CC of Enterococcus faecalis in samples. They can also be used for
CC diagnosing Enterococcal infection in an animal and monitoring
CC progression of disease, and for identifying agents which can be used to
CC modulate the growth or pathogenicity of Enterococcus faecalis, or
CC another related organism, in vivo or in vitro. In particular the
CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC can be used in vaccines to prevent or attenuate an Enterococcal
CC infection.
```

seq_name: N_Geneseq_36:T41700

seq_documentation_block:
 ID T41700 standard; DNA; 1347 BP.
 AC T41700;
 DT 19-JAN-1997 (first entry)
 DE Brugia pahangi beta-tubulin cDNA.
 KW Filariasis; nematode; parasite; beta-tubulin; immunogen; vaccine;
 KW ss.
 OS Brugia pahangi.
 PN WO9632132-A1.
 PD 17-OCT-1996.
 PF 10-APR-1996; U04838.
 PR 10-APR-1995; US-420982.
 PA (UYMC-) UNIV MCGILL.
 PA (UPJO) UPJOHN CO.
 PI Buglio N, Faubert GM, Geary T, Prichard RK;
 DR WPI: 96-476844/47.
 DR P-PSDB: R99425.
 PT New vaccines for filarial parasite infection(s) - comprising
 PT C-terminal beta-tubulin amino acid sequence from a parasite
 PS Example 6; Page 51-53; 67pp; English.
 CC A cDNA clone (T41700) codes for the beta-tubulin (R99425) of
 CC the filarial nematode Brugia pahangi. The C-terminal portion
 CC (see also R99420) of the beta-tubulin is useful in novel vaccines
 CC against filarial parasite infections.
 SQ Sequence 1347 BP; 381 A; 262 C; 337 G; 367 T;

alignment_scores:
 Quality: 32.00 Length: 10
 Ratio: 4.000 Gaps: 0
 Percent Similarity: 80.000 Percent Identity: 70.000

alignment_block:

US-08-653-294-19 x T41700 ..

Align seg 1/1 to: T41700 from: 1 to: 1347

1 TyrArgLeuAlaIle***ArgIleAlaLeu 10
 ||||| ||| |||||:
 998 TACAGATTAGAAATTCATCGTATTCTGTG 1027

seq_name: N_Geneseq_36:Q12225

seq_documentation_block:
 ID Q12225 standard; DNA; 1371 BP.
 AC Q12225;
 DT 13-SEP-1991 (first entry)
 DE Octopus rhodopsin membrane protein.
 KW Octopus; rhodopsin; membrane; helix; OR; ss.
 FH Key Location/Qualifiers
 FT 1..1368
 FT /*tag= a
 FT /*product= membrane protein
 FT misc_feature 322..327
 FT /*tag= a
 FT /*label= SphI
 FT /*note= "restriction enzyme site"
 FT misc_feature 592..597
 FT /*tag= b
 FT /*label= MluI
 FT /*note= "restriction enzyme site"
 FT misc_feature 884..889
 FT /*tag= c
 FT /*label= NdeI
 FT /*note= "restriction enzyme site"
 FT misc_feature 1069..1074
 FT /*tag= d
 FT /*label= NheI
 FT /*note= "restriction enzyme site"
 FT misc_feature 1371
 FT /*tag= e
 FT /*label= BamHI

FT J03123486-A.
 PN 27-MAY-1991.
 PF 06-OCT-1989; 260261.
 PR 06-OCT-1989; JP-260261.
 PA (HITA) HITACHI KK.
 DR WPI: 91-197925/27.
 DR P-PSDB: R12362.
 PT Genetic engineering of membrane protein - by division of protein
 PT into cartridge genes corresp. to helix structure polypeptide(s)
 PS Disclosure; Fig 1; 17pp; Japanese.
 CC A gene cassette is prepd. by division of the OR membrane protein
 CC encoding helix structure polypeptides. Restriction sites are indicated.
 CC A base sequence contg. such a sequence downstream to the tryptophan
 CC regulating gene derived from the E. coli tryptophan operon, the trpL,
 CC or the trpE polypeptide translation regulating base sequence.
 CC and the N-terminal Met of the trpL (or trpE) or trpE polypeptide,
 CC respectively, is introduced in an expression vector for
 CC transformation of host cells.
 SQ Sequence 1371 BP; 298 A; 368 C; 351 G; 354 T;

alignment_scores:
 Quality: 32.00 Length: 10
 Ratio: 3.556 Gaps: 0
 Percent Similarity: 90.000 Percent Identity: 60.000

alignment_block:

US-08-653-294-19 x Q12225/rev ..

Align seg 1/1 to reverse of: Q12225 from: 1 to: 1371

2 ArgLeuAlaIle***ArgIleAlaLeuArg 11
 |||:||||: |||:|||||
 1272 CGGCTAGCCCTCGCGGGTAGCCCTGAGG 1243

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Strd	Orig	zScore	EScore	Len	Documentation
	Sequence				
+	gb_gss8:AQ058120	38.00	144.64	27.25	468
+	gb_gss8:AQ059507	38.00	144.50	27.78	476
+	gb_est44:AV395907	38.00	147.42	27.78	476
+	gb_est20:AA0897834	38.00	142.83	34.40	575
-	gb_gss3:BO0205	37.00	145.58	24.17	263
+	gb_est5:H72837	37.00	139.84	50.47	504
+	gb_gss5:AQ081582	37.00	137.98	64.04	622
+	gb_gss6:AQ085162	37.00	136.64	76.05	724
+	gb_gss6:AQ0864396	37.00	135.83	84.42	794
+	gb_est12:AA320981	36.00	141.03	43.30	275
+	gb_gss1:CNS000WT6	36.00	135.87	84.02	494
+	gb_est24:AI235339	36.00	135.72	85.56	502
+	gb_gss10:AQ0210986	36.00	135.72	85.56	502
+	gb_gss9:AQ0212954	36.00	135.62	86.72	508
+	gb_est27:AI407553	36.00	135.57	87.30	511
+	gb_est44:AA179172	36.00	135.41	89.05	520
+	gb_est10:AA194167	36.00	133.83	109.06	622
+	gb_est35:AI828638	36.00	131.82	140.70	779
+	gb_est35:AI864155	35.00	138.12	62.91	239
+	gb_est41:AW159602	35.00	137.11	71.83	268
+	gb_est11:AQ027930	35.00	135.88	83.83	308
-	gb_est1:AO4318	35.00	133.58	112.69	400
+	gb_est8:AA003505	35.00	132.88	123.26	433
-	gb_est4:R65508	35.00	132.70	126.17	442
+	gb_est21:AA951497	35.00	132.68	126.49	443
+	gb_est38:AL120179	35.00	132.31	132.65	462
-	gb_gss13:AQ038867	35.00	131.75	142.44	492
+	gb_est19:AA0802027	35.00	131.59	145.39	501
+	gb_gss13:AA0392979	35.00	131.38	149.34	513
+	gb_gss4:AQ0677348	35.00	131.21	152.64	523
+	gb_gss10:AQ0225567	35.00	131.21	152.64	523
+	gb_est14:AA439960	35.00	130.94	157.93	539
+	gb_est32:AI729220	35.00	130.73	162.25	552
+	gb_gss15:AQ0581579	35.00	130.37	169.92	575
-	gb_gss10:AQ0569045	35.00	130.06	176.96	596
-	gb_est40:AW142967	35.00	129.87	181.34	609
+	gb_est32:AI726854	35.00	129.72	184.71	619
+	gb_gss15:AQ0662341	35.00	129.36	193.51	645
-	gb_gss6:AQ0840156	35.00	128.88	205.78	681
-	gb_est25:AI256829	35.00	128.30	221.58	727
+	gb_gss11:AQ0327491	35.00	127.49	245.88	797
+	gb_gss4:AQ029455	35.00	127.39	249.02	806
-	gb_est26:AA005267	35.00	127.35	250.42	810

```

KEYWORDS
SOURCE Chlamydomonas reinhardtii.
ORGANISM Chlamydomonas reinhardtii
Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Volvocales;
Chlamydomonadaceae; Chlamydomonas.
REFERENCE
1 (bases 1 to 476)
AUTHORS Asamizu,E., Nakamura,Y., Sato,S., Fukuzawa,H. and Tabata,S.
TITLE A Large Scale Structural Analysis of cDNAs in a Unicellular Green
Alga, Chlamydomonas reinhardtii. I. Generation of 3451
non-redundant Expressed Sequence Tags
JOURNAL DNA Res. (1998) In press
COMMENT On Jun 5, 1998 this sequence version replaced gi:3189716.
Contact: Yasukazu Nakamura
The First Laboratory for Plant Gene Research
Kazusa DNA Research Institute
Yana 1532-3, Kisarazu, Chiba 292-0812, Japan
Email: ynakamu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.
FEATURES
source
1..476
Location/Qualifiers
/organism="Chlamydomonas reinhardtii"
/strain="C9"
/db_xref="taxon:3055"
/clone="CL51a05_r"
/clone_lib="Chlamydomonas reinhardtii C9"
/dev_stage="photoautotrophic growth"
/note="Vector: pBluescriptII SK-; Site_1: EcoRI; Site_2:
XhoI"
BASE COUNT 107 a 124 c 150 g 95 t
ORIGIN
alignment_scores
Quality: 38.00 Length: 10
Ratio: 4.22 Gaps: 0
Percent Similarity: 90.000 Percent Identity: 80.000
alignment_block
US-08-653-294-19 x AV395907 ..
Align seg 1/1 to: AV395907 from: 1 to: 476
3 LeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||:
72 CTGGCCCTGGTACCTATCGCGCTCAGGTAT 101
seq_name: gb_est20:AA897834
seq_documentation_block: 575 bp mRNA EST 12-APR-1998
LOCUS AA897834
DEFINITION NCP3E17 Perithecial Neurospora crassa cDNA clone NP5E1 3' end,
mRNA sequence.
ACCESSION AA897834
VERSION AA897834.1 GI:3044267
KEYWORDS EST.
SOURCE Neurospora crassa.
ORGANISM Neurospora crassa
Eukaryota; Fungi; Ascomycota; Pyrenomycetes; Sordariales;
Sordariaceae; Neurospora.
REFERENCE
1 (bases 1 to 575)
AUTHORS Nelson,M.A., Kang,S., Braun,E.L., Crawford,M.E., Dolan,P.L.,
Leonard,P.M., Mitchell,J., Armijo,A.M., Bean,L., Blueyes,E.,
Cushing,T., Errett,A., Fleharty,M., Gorman,M., Judson,K.,
Miller,R., Ortega,J., Pavlova,I., Perea,J., Todisco,S.,
Trujillo,R., Valentine,J., Wells,A., Werner-Washburne,M., Yazzie,S.
and Natvig,D.O.
Expressed sequences from conidial, mycelial, and sexual stages of
Neurospora crassa
Fungal Genet. Biol. 21, 348-363 (1997)
97435549
COMMENT On Jan 19, 1998 this sequence version replaced gi:2151810.
Contact: Natvig,D.O./Nelson,M.A.
Department of Biology
University of New Mexico
Castetter Hall, Albuquerque, NM 87131, USA
Tel: 505 277 3411
Fax: 505 277 0304
Email: ngp@biology.unm.edu
Deposited in GenBank at the National Center for Genome Resources with
accession GSD8:S:1146747
Seq primer: T7.
FEATURES
source
1..575
Location/Qualifiers
/organism="Neurospora crassa"
/strain="fl a"
/db_xref="taxon:5141"
/clone="NP5E1"
/clone_lib="Perithecial"
/sex="Mating type a (fluffy), fertilized"
/tissue_type="Perithecia (fruiting bodies)"
/dev_stage="perithecia"
/note="mRNA isolated from 5 day old perithecia (fruiting
bodies) of the fluffy strain fl a (Mating type a),
fertilized with conidia from 74-OR23-IV A (Mating type A).
cDNA directionally cloned into pBluescript SK(-) using
the Uni-ZAP XR vector system (Stratagene, La Jolla, CA)."
BASE COUNT 189 a 115 c 90 g 181 t
ORIGIN
alignment_scores
Quality: 38.00 Length: 12
Ratio: 3.800 Gaps: 0
Percent Similarity: 83.333 Percent Identity: 66.667
alignment_block
US-08-653-294-19 x AA897834 ..
Align seg 1/1 to: AA897834 from: 1 to: 575
1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||:|||||:|||||:|||||:|||||:
102 TACAGACTGGATATATCTAAAGTCAAGTTAAGGTAT 137
seq_name: gb_gss3:B02055
seq_documentation_block: 263 bp DNA GSS 13-JUL-1996
LOCUS B02055
DEFINITION CSRL-147D11-u CSRL flow sorted Chromosome 11 specific cosmid Homo
sapiens genomic clone CSRL-147D11, genomic survey sequence.
ACCESSION B02055
VERSION B02055.1 GI:1411333
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 263)
AUTHORS Evans,G.A., Burbee,D., Davies,C., Hahner,L., Oliver,T., Gilbert,M.,
Jones,D., Ward,T., Gillilan,E., Schagemann,J., Probst,S.,
Harris,J., Deford,J., McFarland,J., Burzinski,K., Khan,M.,
Kupfer,K. and Garner,H.R.
Genomic Sequence Sampled Map of Chromosome 11
Unpublished (1996)
CONTACT: Evans GA, Shane Probst
McDermott Center for Human Growth and Development
University of Texas Southwestern Medical Center At Dallas
5323 Harry Hines Blvd, Dallas TX 75235-8591
Tel: 214-648-1600
Fax: 214-648-1666
Email: gevas@utsw.swmed.edu, shane@mcdermott.swmed.edu
PCR Primers
FORWARD: TCTTGAATTTAGGAGTTGTC
BACKWARD: GTTATGTCACCACTATTC
Seq primer: T7
Class: cosmid ends
High quality sequence stop: 263.

```



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source
1..622
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate-924 Col-12 Row-I"
/clone_lib="RPC1-11 Human Male BAC Library"
/sex="male"
/note="vector: pBACE3.6; Genomic sequence of BAC ends"
BASE COUNT 190 a 145 c 123 g 144 t 20 others
ORIGIN

alignment_scores:
  Quality: 37.00      Length: 12
  Ratio: 4.111      Gaps: 0
Percent Similarity: 75.000 Percent Identity: 66.667

alignment_block:
US-08-653-294-19 x A0815582 ..
Align seg 1/1 to: A0815582 from: 1 to: 622

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
||||| ||||| :||||| ||||| |||||
321 TATAGACGCCCATAGATAAGTGGCTCTAAGATAC 356

seq_name: gb_gss6:AQ851612

seq_documentation_block:
LOCUS AQ851612 724 bp DNA GSS 18-OCT-1999
DEFINITION Cpg1352B Cp10WAGNAL Cryptosporidium parvum genomic similar to SKB1
homologue (negative regulator of mitosis) (regulator of Shk1, a
p21(Cdc42/Rac)-activated kinase (PAK)), genomic survey sequence.
ACCESSION AQ851612
VERSION AQ851612.1 GI:6063307
KEYWORDS GSS.
SOURCE Cryptosporidium parvum.
ORGANISM Cryptosporidium parvum.
Eukaryota; Alveolata; Apicomplexa; Coccidia; Eimeriida;
Cryptosporididae; Cryptosporidium.
1 (bases 1 to 724)
Strong, W.B. and Nelson, R.G.
Cryptosporidium parvum GSS Project
Unpublished (1997)
On Sep 10, 1998 this sequence version replaced gi:3553959.
Contact: Nelson, R. G.
Depts. of Medicine & Pharmaceutical Chemistry
San Francisco General Hospital-University of California, San
Francisco
Box 0811, San Francisco, CA 94143-0811, USA
Tel: 415 206 8846
Fax: 415 206 3353
Email: malaria@itsa.ucsf.edu
For Annotation data see http://medsfgh.ucsf.edu/id/cpfrag/home.html
Seq primer: T3
Class: shotgun.
Location/Qualifiers
1..724
/organism="Cryptosporidium parvum"
/strain="IOWA"
/db_xref="taxon:5807"
/clone_lib="Cp10WAGNAL"
/lab_host="E. coli XL2 Blue MRF'"
/note="vector: pBlueScript II (SK-); Site_1: EcoRV; C.
parvum (IOWA isolate) genomic DNA was hydrodynamically
sheared to produce fragments having a tight size
distribution between 2-4 kb by Dr. Yvonne Thorstenson of
the Stanford DNA Sequencing and Technology Center
(http://sequence-
www.stanford.edu/group/techdev/shear.htm). The randomly
sheared gDNA was chromatographed on Sephacryl S-400 to
remove any small fragments and DNA eluting in the void
volume was subcloned into an EcoR V-digested, alkaline
phosphatase-treated pBlueScript II (SK-) vector and

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transformed into E. coli strain XL2 Blue MRF'.
Recombinant clones from the first plating of the library
were selected for sequence analysis using T3 and T7
primers."
BASE COUNT 264 a 93 c 121 g 244 t 2 others
ORIGIN

alignment_scores:
  Quality: 37.00      Length: 11
  Ratio: 4.111      Gaps: 0
Percent Similarity: 81.818 Percent Identity: 81.818

alignment_block:
US-08-653-294-19 x AQ851612/rev ..
Align seg 1/1 to reverse of: AQ851612 from: 1 to: 724

1 TyrArgLeuAlaIle***ArgIleAlaLeuArg 11
||||| ||| :||||| ||||| |||||
552 TATAGCTCTAAATAAGACGTATAGCGTTGAGG 520

seq_name: gb_gss6:AQ864396

seq_documentation_block:
LOCUS AQ864396 794 bp DNA GSS 03-NOV-1999
DEFINITION nbeb0023D13f CUGI Rice BAC Library (ECORI) Oryza sativa genomic
clone nbeb0023D13f, genomic survey sequence.
ACCESSION AQ864396
VERSION AQ864396.1 GI:6214957
KEYWORDS GSS.
SOURCE Oryza sativa.
ORGANISM Oryza sativa
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
euphyllophytes; Spermatophyta; Magnoliophyta; Liliopsida; Poales;
Poaceae; Oryza.
1 (bases 1 to 794)
Wing, R.A. and Dean, R.A.
A BAC End Sequencing Framework to Sequence the Rice Genome
Unpublished (1998)
Contact: Wing, R.A.
Clemson University Genomics Institute
Clemson University
100 Jordan Hall, Clemson, SC 29634, USA
Tel: 864 656 7288
Fax: 864 656 4293
Email: rwing@clemson.edu
Seq primer: TAATACGACCTACTATAGGG
Class: BAC ends
High quality sequence start: 84
High quality sequence stop: 408.
Location/Qualifiers
1..794
/organism="Oryza sativa"
/strain="Japonica"
/cultivar="Nipponbare"
/db_xref="taxon:4530"
/clone="nbeb0023D13f"
/clone_lib="CUGI Rice BAC Library (ECORI)"
/tissue_type="Leaf"
/lab_host="E. coli DH10B"
/note="Vector: pBACindigo; Site_1: EcoRI; Site_2: EcoRI;
Rice is the most important food crop in the world. Half of
the world population, especially those inhabiting highly
populated areas of the humid tropics and subtropics, rely
on rice as their primary source of carbohydrate.
Monocotyledonous rice is a diploid plant (2n=24) with a
haploid genome equivalent of 431 Mbp (Arumuganathan and
Earle, 1991). The relatively small genome of rice, three
times larger than that of Arabidopsis, makes it suitable
for genomic studies. In order to facilitate positional
cloning, physical mapping and genome sequencing of rice,
we have constructed a BAC library from Oryza sativa,

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US-08-653-294-19 x CNS00WT6 ..
Align seg 1/1 to: CNS00WT6 from: 1 to: 494

2 ArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||:||||: :|||||:|||||
416 CGAATGCCAGTAGTCAAGTAGCATGAGGTAC 448

seq_name: gb_est24:AI235399

seq_documentation_block:
LOCUS AI235399 502 bp mRNA EST 31-JAN-1999
DEFINITION EST231961 Normalized rat ovary, Bento Soares Rattus sp. cDNA clone
ROVCR36 3' end, mRNA sequence.
ACCESSION AI235399
VERSION AI235399.1 GI:3828905
KEYWORDS EST.
SOURCE Rattus sp.
ORGANISM Rattus sp.
Eukaryota; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
REFERENCE 1 (bases 1 to 502)
AUTHORS Lee,N.H., Glodek,A., Chandra,I., Mason,T.M., Quackenbush,J.,
Kerlavage,A.R. and Adams,M.D.
Rat Genome Project: Generation of a Rat EST (RESt) Catalog & Rat
Gene Index
Unpublished (1998)
On Jan 19, 1998 this sequence version replaced gi:2151609.
Other ESTs: TC63429
Contact: Lee, NH
ATCC
The Institute for Genomic Research
7112, Medical Center Drive, Rockville, MD 20850, USA
Tel: (301)-838-3529
Fax: (301)-838-0208
Email: nhlee@tigr.org
Seq primer: M13-21.
FEATURES
source
1..502
Location/Qualifiers
/organism="Rattus sp."
/db_xref="ATCC (inhost):2040962"
/db_xref="taxon:10118"
/clone="ROVCR36"
/clone_lib="Normalized rat ovary, Bento Soares"
/notes="Organ: ovary; Vector: pT73Pac; Site_1: EcoRI;
Site_2: NotI"
BASE COUNT 169 a 90 c 85 g 158 t
ORIGIN

alignment_scores:
Quality: 36.00 Length: 12
Ratio: 4.000 Gaps: 0
Percent Similarity: 75.000 Percent Identity: 75.000

alignment_block:
US-08-653-294-19 x AI235399/rev ..
Align seg 1/1 to reverse of: AI235399 from: 1 to: 502

1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
|||||:||||: :|||||:|||||
499 TACAGACTTGCCTACAGCCTTCTATTATGAGGTAT 464

seq_name: gb_gss10:AQ210986

seq_documentation_block:
LOCUS AQ210986 502 bp DNA GSS 18-SEP-1998
DEFINITION HS_3229_A2_D06_MR CIT Approved Human Genomic Sperm Library D Homo
sapiens genomic clone Plate=3229 Col=12 Row=G, genomic survey
sequence.
ACCESSION AQ210986
VERSION AQ210986.1 GI:3619955
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 502)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589

KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 502)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
TITLE Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
JOURNAL Proc. Natl. Acad. Sci. U. S. A. 96 (17), 9739-9744 (1999)
MEDLINE 99380589

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/note="Organ: ovary; Vector: pT7T3pac; Site_1: EcoRI;
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US-08-653-294-19 x AW179172 ..
Align seg 1/1 to: AW179172 from: 1 to: 520
1 TyrArgLeuAlaIle***ArgIleAlaLeuArgTyr 12
:||||| |:|:| |:|:| |:|:| |:|:| |:|:|
459 TTCAGATGCCAGTTGATAGTAGTACGCTAGATAT 494
```

GenCore version 4.5
Copyright (c) 1993 - 2000 Compugen Ltd.

OM protein - protein search, using sw model

Run On: February 8, 2000, 01:29:40 ; Search time 122.56 Seconds
(without alignments)
2.706 Million cell updates/sec

Title: US-08-653-294-20

Perfect score: 63

Sequence: 1 YRLAIRIXRILLY 14

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : A_Geneseq_36.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	42	66.7	12	1 R95429	HLA-B2702 84-79-84
2	42	66.7	12	1 W33798	Peptide B2702.84-7
3	42	66.7	12	1 W33799	Immunomodulating d
4	41	65.1	20	1 R92907	HLA-B2702 CTL modu
5	41	65.1	20	1 R95428	HLA-B2702 84-75-84
6	41	65.1	20	1 W33778	Immunomodulating d
7	37	58.7	469	1 W56793	L. lactis F1 porti
8	36	57.1	20	1 R92909	HLA-B2702 CTL modu
9	36	57.1	20	1 R92908	HLA-B2702 CTL modu
10	36	57.1	20	1 W33791	Peptide B2702.84-7
11	36	57.1	20	1 W33792	Peptide B2702.84-7
12	34	54.0	20	1 R95430	HLA-B2702 84-75/77
13	34	54.0	334	1 R95941	Canine Y5 receptor
14	34	54.0	334	1 W37094	Canis domesticus Y
15	34	54.0	445	1 W15232	Rat neuropeptide Y
16	34	54.0	445	1 W15230	Human neuropeptide Y
17	34	54.0	445	1 W27602	Rat neuropeptide Y
18	34	54.0	445	1 W27604	Human neuropeptide Y
19	34	54.0	445	1 W27603	Rat neuropeptide Y
20	34	54.0	455	1 R95939	Human Y5 receptor
21	34	54.0	455	1 W29447	Human hippocampal
22	34	54.0	455	1 W29413	Human hippocampal
23	34	54.0	455	1 W37093	Homo sapiens hippo
24	34	54.0	456	1 R95940	Rat Y5 receptor. M
25	34	54.0	456	1 W29446	Rat hypothalamic n
26	34	54.0	456	1 W29412	Rat hypothalamic n
27	34	54.0	456	1 W37095	Canis domesticus Y
28	34	54.0	456	1 W37092	Rattus norvegicus
29	34	54.0	466	1 W15233	Mouse neuropeptide
30	34	54.0	893	1 W13899	Thermotoga neapoli
31	34	54.0	893	1 W13900	Thermotoga neapoli
32	34	54.0	893	1 W13901	Thermotoga neapoli
33	34	54.0	893	1 W13902	Thermotoga neapoli
34	34	54.0	893	1 W53918	Wild type Tne poly

35 34 54.0 893 1 W53919 Mutant Tne polymerase
36 34 54.0 893 1 W78758 Tne DNA polymerase
37 34 54.0 893 1 W83974 Tne DNA polymerase
38 34 54.0 893 1 W83973 Tne DNA polymerase
39 34 54.0 893 1 W83980 Tne DNA polymerase
40 34 54.0 893 1 W83981 Tne DNA polymerase
41 34 54.0 893 1 W83982 Tne DNA polymerase
42 34 54.0 893 1 W83983 Tne DNA polymerase
43 34 54.0 893 1 W83984 Tne DNA polymerase
44 34 54.0 893 1 W83985 Tne DNA polymerase
45 34 54.0 893 1 W83976 Tne DNA polymerase

ALIGNMENTS

RESULT 1

R95429 ID R95429 standard; peptide; 12 AA.
AC R95429;
DT 12-NOV-1996 (first entry)
DE HLA-B2702 84-79-84 palindrome.
KW HLA: p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN WO9513288-A1.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LEILAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 95-194027/25.
PT Compos. comprising lymphoid surface membrane proteins - which may
PT inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 12; 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
CC human-leucocyte-associated antigens. This sequence represents the
CC HLA-B2702 84-79-84 palindrome. These sequences can be used to isolate
CC the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
CC protein associated with T-cell activation in mammalian T-cells, and is
CC also immunologically cross reactive with the heat shock protein Hsc70.
CC p74 is found in a limited number of cell types, but is particularly
CC expressed on B and T cells. p74 can be isolated by lysis of a suitable
CC cell with an amphoteric detergent, and then passed through an affinity
CC column containing a covalently bound HLA-B2702 palindromic peptide.
CC Compositions comprising the extracellular fragment of p74 combined with
CC HLA-B2702.80-84 (see R95416), induces calcium influx, and inhibits
CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
CC compounds can be screened for their effect on the cytolytic activity of
CC T-cells, by combining them with the extracellular portion of p74 and
CC determining the amount of binding between the candidate compound and p74.
CC Modulation of CTL activity can be inhibited in a cellular composition
CC containing T-cells and antigen presenting cells (APCs), by adding to the
CC mix the extracellular portion of p74, in an amount sufficient to compete
CC with p74 for the binding of the p74 ligand.
SQ Sequence 12 AA:

Query Match 66.7%; Score 42; DB 1; Length 12;
Best Local Similarity 78.6%; Pred. No. 0.045;
Matches 11; Conservative 0; Mismatches 1; Indels 2; Gaps 1;

OY 1 YRLAIRIXRILLY 14

Db 1 YRLAIR--RIALRY 12

RESULT 2

W33798 ID W33798 standard; peptide; 12 AA.
AC W33798;

19-JUN-1998 (first entry)
 DE Peptide B2702.84-79/79-84 tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha1 domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes, be
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 SQ Sequence 12 AA;

Query Match 56.7%; Score 42; DB 1; Length 12;
 Best Local Similarity 78.6%; Pred. No. 0.045;
 Matches 11; Conservative 0; Mismatches 1; Indels 2; Gaps 1;

QY 1 YRLAIRIXRLLRY 14
 ||||| ||||
 Db 1 YRLAIR--RLLRY 12

RESULT 3
 W33799
 ID W33799 standard; peptide: 12 AA.
 AC W33799;
 DT 19-JUN-1998 (first entry)
 DE Immunomodulating dimer peptide #3.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 PT alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Claim 17; Page 35; 41pp; English.
 CC This sequence represents a specifically claimed immunomodulating
 CC dimer peptide of the invention. A peptide-type compound or variant is

CC claimed which has immunomodulating activity, including the N-terminal
 CC acylated and/or C-terminal amidated or esterified forms of up to 60
 CC amino acids, where the peptide-type compound comprises the formula: A-B,
 CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
 CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
 CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
 CC represents amino acid. The sequence in the brackets may optionally be
 CC absent or truncated at any peptide type bond within the brackets. The
 CC compounds comprise amino acid sequences related to a Class I HLA-B
 CC alpha1 domain (positions 79-84). They can be used to inhibit cytotoxic
 CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
 CC vitro. They can also be used in combination with antigenic peptides or
 CC proteins of interest to activate CTLs. They can also inhibit the
 CC proliferation of T cells in response to anti-CD3. The peptide can be
 CC used for preventing rejection of transplants or for treating autoimmune
 CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
 CC The products can also be used for detection and diagnosis.
 SQ Sequence 12 AA;

Query Match 66.7%; Score 42; DB 1; Length 12;
 Best Local Similarity 78.6%; Pred. No. 0.045;
 Matches 11; Conservative 0; Mismatches 1; Indels 2; Gaps 1;

QY 1 YRLAIRIXRLLRY 14
 ||||| ||||
 Db 1 YRLAIR--RLLRY 12

RESULT 4
 R92907
 ID R92907 standard; peptide: 20 AA.
 AC R92907;
 DT 16-MAY-1996 (first entry)
 DE HLA-B2702 CTL modulating peptide (B2702.84-75/75-84).
 KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;
 KW immunosuppressant; graft versus host disorder; transplantation; therapy;
 KW class I MHC; HLA-B2702.
 OS Synthetic.
 PN W09526979-A1.
 PD 12-OCT-1995.
 PF 05-APR-1995; U04349.
 PR 05-APR-1994; US-222851.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM, Parham P;
 DR WPI: 95-359582/46.
 PT Extension of acceptance period of transplants from MHC unmatched
 PT donor hosts - using Class I B75-84 MHC antigen of the recipient
 PT host
 PS Example 15; Page 36; 80pp; English.
 CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of
 CC class I major histocompatibility complex (MHC) antigens. This sequence
 CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class
 CC I MHC HLA-B2702. These sequences can be used to extend the period of
 CC acceptance by a recipient of a transplant from an MHC unmatched donor.
 CC The peptides are administered to a patient in conjunction with a
 CC subtherapeutic amount of an immunosuppressant. This is administered to
 CC the patient for a limited period of time (compared to the lifetime
 CC administration for current treatments). The peptides particularly
 CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)
 CC of the patient.
 SQ Sequence 20 AA;

Query Match 65.1%; Score 41; DB 1; Length 20;
 Best Local Similarity 55.0%; Pred. No. 0.11;
 Matches 11; Conservative 1; Mismatches 2; Indels 6; Gaps 1;

QY 1 YRLAIRI-----XRLLRY 14
 ||||| ||||
 Db 1 YRLAIRLNRERLRLRY 20

RESULT 5
R95428
ID R95428 standard; peptide; 20 AA.
AC R95428;
DE HLA-B*2702 84-75-84 palindromic.
KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
KW cytolysis; antigen presenting cell.
OS Synthetic.
PN W09513288-AL.
PD 18-MAY-1995.
PF 10-NOV-1994; U12985.
PR 10-NOV-1993; US-150493.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Clayberger C, Krensky AM;
DR WPI: 95-194027/25.
PT Compsns. comprising lymphoid surface membrane proteins - which may
PT inhibit cytolytic activity and differentiation of CTLs.
PS Example; Page 12: 29pp; English.
CC R95413, and R95415-R95431 represent palindromes and fragments of
CC human-leucocyte-associated antigens. This sequence represents the
CC HLA-B*2702 84-75-84 palindromic. These sequences can be used to isolate
CC the protein p74 from a T-cell lysate. p74 is a T-cell surface membrane
CC protein associated with T-cell activation in mammalian T-cells, and is
CC also immunologically cross reactive with the heat shock protein Hsc70.
CC p74 is found in a limited number of cell types, but is particularly
CC expressed on B and T cells. p74 can be isolated by lysis of a suitable
CC cell with an amphoteric detergent, and then passed through an affinity
CC column containing a covalently bound HLA-B*2702 palindromic peptide.
CC Compositions comprising the extracellular fragment of p74 combined with
CC HLA-B*2702.60-84 (see R95416), induces calcium influx, and inhibits
CC cytotoxic T lymphocyte (CTL) differentiation or cytolysis. Candidate
CC compounds can be screened for their effect on the cytolytic activity of
CC T-cells, by combining them with the extracellular portion of p74 and
CC determining the amount of binding between the candidate compound and p74.
CC Modulation of CTL activity can be inhibited in a cellular composition
CC containing T-cells and antigen presenting cells (APCs), by adding to the
CC mix the extracellular portion of p74, in an amount sufficient to compete
CC with p74 for the binding of the p74 ligand.
CC Sequence 20 AA;

Query Match 65.1%; Score 41; DB 1; Length 20;
Best Local Similarity 55.0%; Pred. No. 0.11;
Matches 11; Conservative 1; Mismatches 2; Indels 6; Gaps 1;

Qy 1 YRLAIRI-----XRILLRY 14
|||||: |||||
Db 1 YRLAIRLNRERENLRALRY 20

RESULT 6
W33778
ID W33778 standard; peptide; 20 AA.
AC W33778;
DE 19-JUN-1998 (first entry)
KW Immunomodulating dimer peptide #1.
KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
KW rejection.
OS Synthetic.
OS Homo sapiens.
PN W09744351-AL.
PD 27-NOV-1997.
PF 22-MAY-1997; U08689.
PR 24-MAY-1996; US-653294.
PA (STRD) UNIV LELAND STANFORD JUNIOR.
PI Beulow R, Clayberger C, Krensky AM;
DR WPI: 98-086530/08.
PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
PT alpha-1 domain, used for preventing rejection of transplants or

PT treating autoimmune diseases
PS Claim 16; Page 35; 41pp; English.
CC This sequence represents a specifically claimed immunomodulating
CC dimer peptide of the invention. A peptide-type compound or variant is
CC claimed which has immunomodulating activity, including the N-terminal
CC acylated and/or C-terminal amidated or esterified forms of up to 60
CC amino acids, where the peptide-type compound comprises the formula: A-B,
CC where A, B = (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or
CC V; aa77 = D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a
CC hydrophobic or small amino acid; aa82 = R or L; aa83 = G or R; and aa
CC represents amino acid. The sequence in the brackets may optionally be
CC absent or truncated at any peptide type bond within the brackets. The
CC compounds comprise amino acid sequences related to a Class I HLA-B
CC alpha1 domain (positions 79-84). They can be used to inhibit cytotoxic
CC T-lymphocytes (CTL) from undesirably attacking cells in a host or in
CC vitro. They can also be used in combination with antigenic peptides or
CC proteins of interest to activate CTLs. They can also inhibit the
CC proliferation of T cells in response to anti-CD3. The peptide can be
CC used for preventing rejection of transplants or for treating autoimmune
CC diseases, e.g. diabetes, rheumatoid arthritis and lupus erythematosus.
CC The products can also be used for detection and diagnosis.
SQ Sequence 20 AA;

Query Match 65.1%; Score 41; DB 1; Length 20;
Best Local Similarity 55.0%; Pred. No. 0.11;
Matches 11; Conservative 1; Mismatches 2; Indels 6; Gaps 1;

Qy 1 YRLAIRI-----XRILLRY 14
|||||: |||||
Db 1 YRLAIRLNRERENLRALRY 20

RESULT 7

W56793
ID W56793 standard; Protein; 469 AA.
AC W56793;
DE L. lactis F1 portion of F0F1 ATPase beta subunit.
DE Biomass production; uncoupled ATPase; F0F1 ATPase; membrane bound;
KW F1; Lactococcus lactis.
OS Lactococcus lactis.
PN W09810089-AL.
PD 12-MAR-1998.
PF 08-SEP-1997; DK0373.
PR 06-SEP-1996; DK-000963.
PA (JENS/) JENSEN P R.
PI Snoep JL, Westerhoff HV;
DR WPI: 98-193637/17.
DR N-PSDB; V29571.
PT Method improving production of biomass or a desired product - by
PT expressing an uncoupled ATPase activity in the cell
PS Example 5; Pages 45-46; 78pp; English.
CC This is the beta subunit of soluble part (F1) of membrane bound (F0F1
CC type) H⁺-ATPase. The DNA encoding this or a portion of F1 exhibiting
CC ATPase activity is derived from Lactococcus lactis subsp. cremoris strain
CC MG1363 and is used in a novel method for improving the production of
CC biomass or a desired product from a cell. The method comprises expressing
CC an uncoupled ATPase activity in the cell to induce conversion of ATP to
CC ADP without primary effects on other cellular metabolites or functions
CC and incubating the cell with a suitable substrate to produce the biomass
CC or product. The expression is directed using a vector including DNA
CC encoding the soluble part (F1) of the membrane bound (F0F1 type)
CC H⁺-ATPase or a portion of F1 exhibiting ATPase activity, the DNA being
CC derived from Lactococcus lactis subsp. cremoris, Lactococcus lactis
CC subsp. lactis, Streptococcus thermophilus, Phaffia rhodozyma or
CC Trichoderma reesei, where the DNA is under the control of a promoter. An
CC ideal ATPase is the membrane bound H⁺ATPase. This enzyme complex
CC consists of two parts, the membrane integral part and the (F0) and the
CC cytoplasmic part (F1). Together the two parts couple the hydrolysis of
CC ATP and ADP to the translocation of protons across the cytoplasmic
CC membrane, or vice versa. The proton gradient is used to drive ATP
CC synthesis from ADP and P_i. The method can be used for optimising the

Query Match 57.1%; Score 36; DB 1; Length 20;
 Best Local Similarity 50.0%; Pred. No. 0.95;
 Matches 10; Conservative 1; Mismatches 3; Indels 6; Gaps 1;

Oy 1 YRLAIRI-----XRILLRY 14
 ||||| |
 Db 1 YRLATRLNRERENLRALRY 20

RESULT 11

W33792
 ID W33792 standard; peptide; 20 AA.
 AC W33792;
 DT 19-JUN-1998 (first entry)
 DE Peptide B2702 84-75/75-84T tested for immunomodulating activity.
 KW Immunomodulating dimer; immunosuppressant drug; CTL activation;
 KW transplantation; autoimmune disease; Class I HLA-B alpha-1 domain;
 KW rejection.
 OS Synthetic.
 OS Homo sapiens.
 PN W09744351-A1.
 PD 27-NOV-1997.
 PF 22-MAY-1997; U08689.
 PR 24-MAY-1996; US-653294.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Beulow R, Clayberger C, Krensky AM;
 DR WPI: 98-086530/08.
 PT New immunomodulating dimer peptide(s) - based on a Class I HLA-B
 alpha-1 domain, used for preventing rejection of transplants or
 PT treating autoimmune diseases
 PS Example 1; Page 19; 41pp; English.
 CC Peptides W33784-98 and W33778-9 were assayed for their immunomodulating
 CC activity. A peptide-type compound or variant is claimed which has
 CC immunomodulating activity, including the N-terminal acylated and/or
 CC C-terminal amidated or esterified forms of up to 60 amino acids, where
 CC the peptide-type compound comprises the formula: A-B, where A, B =
 CC (R aa76-77L) (aa79-84) or (aa84-79) (Laa77-76R); aa76 = E or V; aa77 =
 CC D, S or N; aa79 = R or G; aa80 = I or N; aa81, aa84 = a hydrophobic or
 CC small amino acid; aa82 = R or L; aa83 = G or R; and aa represents amino
 CC acid. The sequence in the brackets may optionally be absent or truncated
 CC at any peptide type bond within the brackets. The compounds comprise
 CC amino acid sequences related to a Class I HLA-B alpha1 domain (positions
 CC 79-84). They can be used to inhibit cytotoxic T-lymphocytes (CTL) from
 CC undesirably attacking cells in a host or in vitro. They can also be
 CC used in combination with antigenic peptides or proteins of interest to
 CC activate CTLs. They can also inhibit the proliferation of T cells in
 CC response to anti-CD3. The peptide can be used for preventing rejection
 CC of transplants or for treating autoimmune diseases, e.g. diabetes,
 CC rheumatoid arthritis and lupus erythematosus. The products can also be
 CC used for detection and diagnosis.
 CC Sequence 20 AA;

Query Match 57.1%; Score 36; DB 1; Length 20;
 Best Local Similarity 50.0%; Pred. No. 0.95;
 Matches 10; Conservative 1; Mismatches 3; Indels 6; Gaps 1;

Oy 1 YRLAIRI-----XRILLRY 14
 ||||| |
 Db 1 YRLAIRLNRERENLRALRY 20

RESULT 12

R95430
 ID R95430 standard; peptide; 20 AA.
 AC R95430;
 DT 12-NOV-1996 (first entry)
 DE HLA-B2702 84-75/75-84T palindromic.
 KW HLA; p74; alpha1-helix; human-leucocyte-associated antigen; inhibitor;
 KW T-cell lysate; membrane protein; mammal; heat shock protein; Hsc70; APC;
 KW B cell; calcium influx; cytotoxic T lymphocyte; CTL; differentiation;
 KW cytolysis; antigen presenting cell.

OS Synthetic.
 PN W09513288-A1.
 PD 18-MAY-1995.
 PF 10-NOV-1994; U12985.
 PR 10-NOV-1993; US-150493.
 PA (STRD) UNIV LELAND STANFORD JUNIOR.
 PI Clayberger C, Krensky AM;
 DR WPI: 95-194027/25.
 PT Compsns. comprising lymphoid surface membrane proteins - which may
 PT inhibit cytolytic activity and differentiation of CTLs.
 PS Example; Page 12; 29pp; English.
 CC R95413, and R95415-R95431 represent palindromes and fragments of
 CC human-leucocyte-associated antigens. These sequences can be used to
 CC HLA-B2702 84-75/75-84T palindromic.
 CC isolate the protein p74 from a T-cell lysate. p74 is a T-cell surface
 CC membrane protein associated with T-cell activation in mammalian T-cells,
 CC and is also immunologically cross reactive with the heat shock protein
 CC Hsc70. p74 is found in a limited number of cell types, but is
 CC particularly expressed on B and T cells. p74 can be isolated by lysis of
 CC a suitable cell with an amphoteric detergent, and then passed through an
 CC affinity column containing a covalently bound HLA-B2702 palindromic
 CC peptide. Compositions comprising the extracellular fragment of p74
 CC combined with HLA-B2702.60-84 (see R95416), induces calcium influx, and
 CC inhibits cytotoxic T lymphocyte (CTL) differentiation or cytolysis.
 CC candidate compounds can be screened for their effect on the cytolytic
 CC activity of T-cells, by combining them with the extracellular portion of
 CC p74 and determining the amount of binding between the candidate compound
 CC and p74. Modulation of CTL activity can be inhibited in a cellular
 CC composition containing T-cells and antigen presenting cells (APCs), by
 CC adding to the mix the extracellular portion of p74, in an amount
 CC sufficient to compete with p74 for the binding of the p74 ligand.
 CC Sequence 20 AA;

Query Match 54.0%; Score 34; DB 1; Length 20;
 Best Local Similarity 52.6%; Pred. No. 2.2;
 Matches 10; Conservative 1; Mismatches 2; Indels 6; Gaps 1;

Oy 1 YRLAIRI-----XRILLR 13
 ||||| |
 Db 1 YRLAIRLNRERENLRALR 19

RESULT 13

R95941
 ID R95941 standard; Protein; 334 AA.
 AC R95941;
 DT 14-OCT-1996 (first entry)
 DE Canine Y5 receptor.
 KW Y5 receptor; atypical neuropeptide Y1 receptor; feeding behavior;
 KW G protein-coupled receptor; agonist; antagonist; obesity;
 KW bulimia; anorexia.
 OS Canis familiaris.
 PN W09616542-A1.
 PD 06-JUN-1996.
 PF 01-DEC-1995; U15646.
 PR 02-DEC-1994; US-349025.
 PA (SYNA) SYNAPTIC PHARM CORP.
 PI Branchek T, Gerald CPG, Walker MW, Weinshank RL;
 DR WPI: 96-277371/28.
 DR N-PSDB; T30435.
 PT Modifying feeding behaviour using Y5 receptor (ant)agonists -
 PT increases or decreases food consumption, for treatment of e.g.
 PT obesity or bulimia
 PS Claim 58; Fig 15; 235pp; English.
 CC Canine Y5 receptor (R95941) was identified as the homologue of rat
 CC hypothalamic Y5 receptor (R95940), isolated as an 'atypical Y1
 CC receptor'. The receptor belongs to the G protein-coupled receptor
 CC superfamily. It is encoded by a cDNA clone (see also T30435) that
 CC was isolated by PCR amplification using primers (T30436-37) based
 CC on human and rat cDNA clones (T30433-34). Recombinant canine Y5
 CC receptor can be produced in prokaryotic or eukaryotic (e.g. COS,
 CC 293 or Sf9 insect) host cells. It is used to identify Y5 ligands

CC (agonists and antagonists) that can be used to treat obesity,
 CC bulimia or anorexia.
 SQ Sequence 334 AA;

Query Match 54.0%; Score 34; DB 1; Length 334;
 Best Local Similarity 42.9%; Pred. No. 34;
 Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
 ||:| | :|:|
 Db 95 YRIAFITISLLVQY 108

RESULT 14

W37094
 ID W37094 standard; Protein; 334 AA.

AC W37094;
 DT 08-JUN-1998 (first entry)
 DE Canis domesticus Y5 receptor.
 DE Y5 receptor; treatment; anorexia; bulimia; obesity;
 KW feeding behaviour; modification; atypical neuropeptide.
 OS Canis domesticus.
 OS W09746250-AA.

PD 11-DEC-1997.
 PF 04-JUN-1997; U09504.
 PR 21-FEB-1997; US-803600.
 PR 04-JUN-1996; US-668650.
 PA (SYNA-) SYNAPTIC PHARM CORP.
 PI Branchek T, Gerald CP, Walker MW, Weinshank RL;
 DR WPI; 98-051901/05.
 DR N-PSDB; V00639.

PT DNA encoding canine hypothalamic atypical neuro:peptide Y/peptide YY
 receptor, Y5 - useful for identification of compounds which are
 PT capable of modifying feeding behaviour
 PS Claim 5: Fig 15: 273pp; English.

CC The sequence is that of a Y5 receptor (Y5-R).
 CC Y5-R can be used in processes to determine whether a chemical compound
 CC specifically binds to and activates or inhibits a Y5-R by measuring a
 CC second messenger response. The chemical compounds can be used to
 CC increase or reduce the activity of a Y5-R. In particular, inhibitors
 CC can be used to treat obesity and activators can be used to treat
 CC anorexia. Antagonists capable of alleviating (by decreasing the
 CC activity of Y5-R) an abnormality can be identified by administering a
 CC potential antagonist to a transgenic mammal as above, and determining
 CC whether the substance alleviates the physical and behavioural
 CC abnormalities displayed by the transgenic mammal as a result of
 CC overactivity of a Y5-R. Agonists can be identified in a similar manner,
 CC but where the abnormality is alleviated by increasing the activity of
 CC Y5-R.

SQ Sequence 334 AA;

Query Match 54.0%; Score 34; DB 1; Length 334;
 Best Local Similarity 42.9%; Pred. No. 34;
 Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
 ||:| | :|:|
 Db 95 YRIAFITISLLVQY 108

RESULT 15

W15232
 ID W15232 standard; Protein; 445 AA.

AC W15232;
 DT 21-JUL-1997 (first entry)
 DE Rat neuropeptide Y-Y5 receptor.
 KW Neuropeptide Y-Y5; appetite; obesity; G-protein coupled receptor;
 KW antidiabetic; hypotensive; neuronal growth factor;
 KW cardiovascular drug; anti-psychotic; neuroleptic; antidiabetic;
 KW agonist; antagonist.
 OS Rattus sp.

PN W09717440-AA.
 PD 15-MAY-1997.
 PF 08-NOV-1996; AU0706.
 PR 09-NOV-1995; AU-006467.
 PA (GARV-) GARVAN INST MEDICAL RES.
 PI Herzog H;
 DR WPI; 97-281029/25.
 DR N-PSDB; T66911.
 DR N-PSDB; T66910.

PT DNA encoding the neuropeptide Y-Y5 receptor - for screening for
 PT NPY-Y5 antagonists and agonists, useful as anti-obesity agents,
 PT anti-hypertensive agents cardiovascular drugs, etc.
 PS Claim 17; Fig 3; 44pp; English.

CC A novel rat neuropeptide Y (NPY)-Y1-like receptor (W15232),
 CC designated NPY-Y5 receptor, is a G-protein coupled receptor of
 CC NPY, which is involved in appetite/obesity regulation. Its amino
 CC acid sequence was deduced from a cDNA clone (T66911) isolated from
 CC a rat hypothalamic library. Human (W15230) and mouse (W15233)
 CC NPY-Y5 receptors have also been identified. The receptors can be
 CC expressed on the cell surface of host (pref. CHO, human embryonic
 CC kidney 293 or insect Sf9) cells. The receptors or host cells can
 CC be used to screen for (ant)agonists of NPY useful as potential
 CC hypotensives, cardiovascular drugs, neuronal growth factors,
 CC anti-psychotic, anti-obesity or anti-diabetic drugs.
 SQ Sequence 445 AA;

Query Match 54.0%; Score 34; DB 1; Length 445;
 Best Local Similarity 42.9%; Pred. No. 45;
 Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
 ||:| | :|:|
 Db 207 YRIAFITISLLVQY 220

Search completed: February 8, 2000, 01:29:41
 Job time: 1753 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:30 ; Search time 117.7 Seconds
(without alignments)
5.611 Million cell updates/sec

Title: US-08-653-294-20

Perfect score: 63

Sequence: 1 YRLAIRIXRILLRY 14

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : PIR_62.*

1: pir1.*

2: pir2.*

3: pir3.*

4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	39	61.9	485	2 T09974	H+-transporting ATP
2	39	61.9	486	2 B70775	probable atpD prot
3	38	60.3	133	2 D72110	hypothetical prote
4	38	60.3	194	2 F64075	urease accessory p
5	38	60.3	478	2 H70474	ATP synthase F1 be
6	38	60.3	709	1 OXCKPM	acyl-CoA oxidase (
7	36	57.1	96	2 B69327	conserved hypother
8	36	57.1	502	1 OXCKAX	acyl-CoA oxidase (
9	36	57.1	709	1 OXCKX4	acyl-CoA oxidase (
10	36	57.1	709	1 OXCKX	acyl-CoA oxidase (
11	36	57.1	3131	2 S39842	emiatin synthetas
12	35	55.6	724	1 OXCKP2	acyl-CoA oxidase (
13	35	55.6	724	2 Jc4563	acyl-CoA oxidase (
14	34	54.0	157	2 F72612	hypothetical prote
15	34	54.0	276	2 F72721	probable citrate 1
16	34	54.0	323	2 A70404	acetyl-CoA carboxy
17	34	54.0	662	1 OXCKX5	acyl-CoA oxidase (
18	34	54.0	2009	2 S49764	SEC7 protein - yea
19	33	52.4	149	2 G72250	ribosomal protein
20	33	52.4	260	2 S57939	Ahrv protein - Aer
21	33	52.4	331	2 G75035	iron (iii) abc tra
22	33	52.4	470	2 S30597	H+-transporting AT
23	33	52.4	478	2 S37547	H+-transporting AT
24	33	52.4	485	2 A70706	probable phor prot
25	33	52.4	532	2 G70986	probable coA ligas
26	33	52.4	1729	2 A49282	fusion protein la/
27	33	52.4	2376	2 S48405	probable membrane
28	33	52.4	2630	2 T08868	polyprotein p1 - A
29	32	50.8	205	2 T10296	fibroblast growth
30	32	50.8	339	2 D72509	hypothetical prote

31 32 50.8 489 2 T13026 hypothetical prote
32 32 50.8 591 2 B72086 hypothetical prote
33 32 50.8 785 2 T01541 hypothetical prote
34 31 49.2 113 1 R5BY1E ribosomal protein
35 31 49.2 113 2 S55962 ribosomal protein
36 31 49.2 185 2 S74416 hypothetical prote
37 31 49.2 195 2 S74416 hypothetical prote
38 31 49.2 220 2 G72472 pituitary adenylat
39 31 49.2 224 2 F71329 hypothetical prote
40 31 49.2 277 2 Jc2565 probable phosphogl
41 31 49.2 294 2 H38888 replication protei
42 31 49.2 331 2 A71128 COI intron 15 prot
43 31 49.2 334 2 G69303 probable iron (iii
44 31 49.2 343 2 S15949 iron (iii) ABC tra
45 31 49.2 351 2 H70570 hypothetical prote

ALIGNMENTS

RESULT 1

T09974

H+-transporting ATP synthase (EC 3.6.1.34) beta chain - Mycobacterium leprae
C:Species: Mycobacterium leprae
C:Date: 20-Sep-1999 #sequence_revision 20-Sep-1999 #text_change 20-Sep-1999
C:Accession: T09974

R:Robison, K.

submitted to the EMBL Data Library, September 1994

A:Reference number: Z16911

A:Accession: T09974

A:Status: translated from GB/EMBL/DBDJ

A:Molecule type: DNA

A:Residues: 1-485 <ROB>

A:Cross-references: EMBL:U15186; NID:g699323; PID:g699347

C:Keywords: ATP biosynthesis; hydrolase; membrane-associated complex

Query Match 61.9%; Score 39; DB 2; Length 485;

Best Local Similarity 57.1%; Pred. No. 3.7;

Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14

DB 376 YRVAQEVIRILQRY 389

RESULT 2

B70775

probable atpD protein - Mycobacterium tuberculosis (strain H37RV)

C:Species: Mycobacterium tuberculosis

C:Date: 17-Jul-1998 #sequence_revision 17-Jul-1998 #text_change 16-Jul-1999

C:Accession: B70775

R:Cole, S.T.; Brosch, R.; Parkhill, J.; Garnier, T.; Churcher, C.; Harris, D.; Gordon

; Connor, R.; Davies, R.; Devlin, K.; Feltwell, T.; Gentles, S.; Hamlin, N.; Holtroyd,

Rajandream, M.A.; Rogers, J.; Rutter, S.; Seeger, K.; Skelton, S.; Squares, S.

Nature 393, 537-544, 1998

A:Authors: Squires, R.; Sulston, J.E.; Taylor, K.; Whitehead, S.; Barrrell, B.G.

A:Title: Deciphering the biology of Mycobacterium tuberculosis from the complete geno

A:Reference number: A70500; MUID:98295987

A:Accession: B70775

A:Status: preliminary; nucleic acid sequence not shown; translation not shown

A:Molecule type: DNA

A:Residues: 1-486 <COI>

A:Cross-references: GB:Z73419; GB:AL123456; NID:g3261573; PIDN:CAA97743.1; PID:e24365

A:Experimental source: strain H37RV

C:Genetics:

A:Gene: atpD

C:Superfamily: H+-transporting ATP synthase alpha chain; H+-transporting ATP synthase

F:197-366/Domain: H+-transporting ATP synthase alpha chain homology <ATP>

Query Match 61.9%; Score 39; DB 2; Length 486;

Best Local Similarity 57.1%; Pred. No. 3.7;

Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
||:| : ||| ||
Db 377 YRVAQEVIRILQRY 390

RESULT 3

D72110
hypothetical protein - Chlamydia pneumoniae (strain CWL029)
C:Species: Chlamydia pneumoniae
C>Date: 23-Apr-1999 #sequence_revision 23-Apr-1999 #text_change 23-Apr-1999
C:Accession: D72110
R:Kallman, S.; Mitchell, W.; Marathe, R.; Lammel, C.; Fan, J.; Olinger, L.; Grimwood, J.;
Nature Genet. 21, 385-389, 1999
A:Title: Comparative genomes of Chlamydia pneumoniae and C. trachomatis.
A:Reference number: A72000; MUID:99206606
A:Accession: D72110
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-133 <ARN>
A:Cross-references: GB:AE001604; GB:AE001363; NID:g4376438; PID:g4376450
A:Experimental source: strain CWL029
C:Genetics:
A:Gene: CPn0181

Query Match 60.3%; Score 38; DB 2; Length 133;

Best Local Similarity 61.5%; Pred. No. 1.7; Indels 0; Gaps 0;

Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 RLAIRIXRILLRY 14
| : | : |||||
Db 88 RIPWRLKIRILLRY 100

RESULT 4

F64075
urase accessory protein homolog - Haemophilus influenzae (strain Rd KW20)
C:Species: Haemophilus influenzae
C>Date: 18-Aug-1995 #sequence_revision 18-Aug-1995 #text_change 31-Oct-1997
C:Accession: F64075
R:Fleischmann, R.D.; Adams, M.D.; White, O.; Clayton, R.A.; Kirkness, E.F.; Kerlavage, A.;
Gocayne, J.D.; Scott, J.; Shirley, R.; Liu, L.I.; Glodek, A.; Kelley, J.M.; Weidman, J.;
D.M.; Brandon, R.C.; Fine, L.D.; Fritchman, J.L.; Fuhrmann, J.L.; Geoghegan, N.S.M.
Science 269, 496-512, 1995
A:Authors: Gnehm, C.L.; McDonald, L.A.; Small, K.V.; Fraser, C.M.; Smith, H.O.; Venter, A.;
A:Title: Whole-genome random sequencing and assembly of Haemophilus influenzae Rd.
A:Reference number: A64000; MUID:95350630
A:Accession: F64075
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-194 <TIGR>
A:Cross-references: GB:U32736; GB:I42023; NID:gl573519; PID:gl573522; TIGR:HI0537
C:Genetics:
A:Start codon: GTG

Query Match 60.3%; Score 38; DB 2; Length 194;

Best Local Similarity 35.7%; Pred. No. 2.4; Indels 0; Gaps 0;

Matches 5; Conservative 6; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
||:| : ||| ||
Db 64 PKLGVRLKIFIRY 77

RESULT 5

H70474
ATP synthase F1 beta subunit - Aquifex aeolicus
C:Species: Aquifex aeolicus
C>Date: 08-May-1998 #sequence_revision 08-May-1998 #text_change 16-Jul-1999
C:Accession: H70474

R;Deckert, G.; Warren, P.V.; Gaasterland, T.; Young, W.G.; Lenox, A.L.; Graham, D.E.;
V.
Nature 392, 353-358, 1998
A:Title: The complete genome of the hyperthermophilic bacterium Aquifex aeolicus.
A:Reference number: A70300; MUID:98196686
A:Accession: H70474
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-478 <QOF>
A:Cross-references: GB:AE000769; NID:g2984262; PIDN:RAC07790.1; PID:g2984264; GB:AE00
A:Experimental source: strain VF5
C:Genetics:
A:Gene: atpD

C:Superfamily: H+-transporting ATP synthase alpha chain; H+-transporting ATP synthase
C:Keywords: P-loop
F;163-170/Region: nucleotide-binding motif A (P-loop)
F;189-358/Domain: H+-transporting ATP synthase alpha chain homology <ATP>

Query Match 60.3%; Score 38; DB 2; Length 478;

Best Local Similarity 50.0%; Pred. No. 5.7;

Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
| : | : |||||
Db 369 YEVAMEVKRILQRY 382

RESULT 6

OXCKPM
acyl-CoA oxidase (EC 1.3.3.6) PXP4, peroxisomal - yeast (Candida maltosa)
C:Species: Candida maltosa
C>Date: 31-Mar-1989 #sequence_revision 31-Mar-1989 #text_change 17-Feb-1995
C:Accession: A29441
R:Hill, D.E.; Boulay, R.; Rogers, D.
Nucleic Acids Res. 16, 365-366, 1988
A:Title: Complete nucleotide sequence of the peroxisomal acyl CoA oxidase from the al
A:Reference number: A29441; MUID:88124223
A:Accession: A29441
A:Molecule type: DNA
A:Residues: 1-709 <HIL>
A:Experimental source: ATCC 20184
C:Genetics:
A:Gene: POX4
C:Superfamily: acyl-CoA oxidase
C:Keywords: FAD; fatty acid oxidation; flavoprotein; oxidoreductase; peroxisome

Query Match 60.3%; Score 38; DB 1; Length 709;

Best Local Similarity 50.0%; Pred. No. 8.4;

Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
||:| : ||| ||
Db 308 YRLARVSTIALRY 321

RESULT 7

B69327
conserved hypothetical protein AF0618 - Archaeoglobus fulgidus
C:Species: Archaeoglobus fulgidus
C>Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 05-Jun-1998
C:Accession: B69327
R;Klenk, H.P.; Clayton, R.A.; Tomb, J.F.; White, O.; Nelson, K.E.; Ketchum, K.A.; Dod
; Fleischmann, R.D.; Quackenbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E
Glodek, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L.
Nature 390, 364-370, 1997
A:Authors: Utterback, T.; Cotton, M.D.; Spriggs, T.; Artiach, P.; Kaine, B.P.; Sykes,
Smith, H.O.; Woese, C.R.; Venter, J.C.
A:Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing arch
A:Reference number: A69250; MUID:98049343
A:Accession: B69327
A:Status: preliminary; nucleic acid sequence not shown; translation not shown

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WASHINGTON, D.C.

Search completed: February 7, 2000, 11:54:32
Job time: 24342 sec

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OM protein - protein search, using sw model

Run on: February 8, 2000, 00:59:58 ; Search time 63.71 Seconds
(without alignments)
6.563 Million cell updates

Title: US-08-653-294-20
Perfect score: 63
Sequence: 1 YRLAIRIXRILLRY

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 82229 seqs, 29864866 residues

Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : SwissProt_38:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query		DB	ID	Description
		Match	Length			
1	42	66.7	227	1	UREF_ACTPL	O54423 actinobacil
2	39	61.9	485	1	ATPB_MYCLE	P45823 mycobacteri
3	39	61.9	486	1	ATPB_MYCTO	Q10593 mycobacteri
4	38	60.3	194	1	UREF_HAEIN	P44395 haemophilus
5	38	60.3	471	1	ATPB_HERAU	P42466 herpetosiph
6	38	60.3	478	1	ATPB_AQUAE	O67828 aquifex aeo
7	38	60.3	478	1	ATPB_AQUPY	O50292 aquifex pyr
8	38	60.3	708	1	CAO4_CANMA	P05335 candida mal
9	36	57.1	502	1	CAO3_CANTR	P11355 candida tro
10	36	57.1	708	1	CAO2_CANTR	P06598 candida tro
11	35	55.6	723	1	CAO4_CANTR	P11356 candida tro
12	35	55.6	724	1	CAO2_CANMA	Q00468 candida mal
13	34	54.0	323	1	ACCA_AQUAE	O00468 candida mal
14	34	54.0	446	1	NY5R_CANFA	O67260 aquifex aeo
15	34	54.0	446	1	NY5R_PIG	O62729 canis famli
16	34	54.0	455	1	NY5R_HUMAN	O97969 sus scrofa
17	34	54.0	456	1	NY5R_RAT	Q15761 homo sapien
18	34	54.0	466	1	NY5R_MOUSE	Q63634 rattus norv
19	34	54.0	466	1	CAO1_CANTR	O70342 mus musculu
20	34	54.0	661	1	CAO1_CANTR	O03790 candida tro
21	33	52.4	260	1	SEC7_YEAST	P11075 saccharomyc
22	33	52.4	260	1	AHYR_AERYH	Q44059 aeromonas h
23	33	52.4	470	1	ATPB_LACCA	Q03234 lactobacill
24	33	52.4	477	1	ATPB_STRLI	P50004 streptomyce
25	33	52.4	2376	1	YIN9_YEAST	P40468 saccharomyc
26	32.5	51.6	2569	1	LM33_MOUSE	Q61789 mus musculu
27	32	50.8	205	1	FGPH_NPVPQ	Q10284 orgyia pseu
28	31	49.2	468	1	ATPB_ENTHR	P43451 enterococcu
29	31	49.2	112	1	RL31_YEAST	P04649 saccharomyc
30	31	49.2	195	1	PACA_CLAMA	P48144 clarias mac
31	31	49.2	343	1	YSC2_THEEL	P25125 thermus aqu
32	31	49.2	626	1	YIT7_MYCTO	Q10966 mycobacteri
33	31	49.2	1603	1	VIT5_CAEEL	P06125 caenorhabdi
34	31	49.2	3433	1	UTRO_HUMAN	P46939 homo sapien
35	30	47.6	57	1	YCU5_CAEEL	O22702 caenorhabdi

	35	30	47.6	72	1	YVAV_VACC	P20530	vaccinia vi
35	35	30	47.6	72	1	YVAV_VACC	P20530	vaccinia vi
36	36	30	47.6	133	1	R86_CHLIR	P28544	chlamydia t
37	30	47.6	142	1	RLI3_PYRHO	O59300	pyrococcus	
38	30	47.6	207	1	YTOR_CVBH	P22854	bovine coro	
39	38	30	47.6	207	1	YTOR_CVBH	P10525	bovine coro
40	39	30	47.6	207	1	YTOR_CVBH	P26626	turkey ente
41	30	47.6	207	1	YTOR_CVKKE	P52133	escherichia	
42	41	30	47.6	233	1	YFJR_ECOLI	Q09465	caenorhabdi
43	41	30	47.6	274	1	O59_CABEL	P77365	escherichia
44	43	30	47.6	285	1	YAFY_ECOLI	P77365	escherichia
45	43	30	47.6	289	1	LIP1_SVNV3	P52980	synechocyst
46	44	30	47.6	326	1	HOLB_PSAE	P72024	pseudomonas

ALIGNMENTS

```

RESULT 1
UREF_ACTPL STANDARD; PRT; 227 AA.
ID ID UREF_ACTPL
AC O54423;
DT 15-DEC-1998 (Rel. 37, Created)
DT 15-DEC-1998 (Rel. 37, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE UREASE ACCESSORY PROTEIN UREF.
DE UREF.
GN Actinobacillus pleuropneumoniae (Haemophilus pleuropneumoniae).
OS Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
OC Actinobacillus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CM5;
RA BOSSE J.T., MACINNES J.I.;
RL Submitted (FEB-1997) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PROBABLY FACILITATING NICKEL INCORPORATION.
CC -1- SIMILARITY: BELONGS TO THE UREF FAMILY.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; U89957; AAC00063.1; -.
DR DR
KW Nickel.
SO SEQUENCE 227 AA; 25397 MW; 310CB946 CRC32;

```

```
Query Match          66.7%; Score 42; DB 1; Length 227;
Best Local Similarity 42.9%; Pred. No. 0.16;
Matches 6: Conservative 5; Mismatches 3; Indels 0; Gaps 0;
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Qy 1 YRLAIRXRLLRY 14
| : | : | : | :
Db 97 YKLGVRLLKFIRY 110

RESULT	2				
ATPB_MYCLE					
ID	ATPB_MYCLE	STANDARD;	PRT;	485	AA.
AC	P45823;				
DT	01-NOV-1995	(Rel. 32; Created)			
DT	01-NOV-1995	(Rel. 32; Last sequence update)			
DT	01-NOV-1995	(Rel. 32; Last annotation update)			
DE	ATP SYNTHASE BETA CHAIN (EC 3.6.1.34).				
GN	ATPD.				

OC *Mycobacterium leprae*.
 CS Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
 [1]
 RN
 RP SEQUENCE FROM N.A.
 RA SMITH D.R., ROBISON K.;

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RL Submitted (SEP-1994) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON
CC GRADIENT ACROSS THE MEMBRANE. THE BETA CHAIN IS THE CATALYTIC
CC SUBUNIT.
CC -!- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC
CC CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
CC HAS THREE MAIN SUBUNITS: A, B AND C.
CC -!- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
CC -----
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CC -----
DR EMBL; U15186; AAA63108.1; -
DR HSSP; P07677; 1SKY.
DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
DR PFAM; PF00006; ATP-synt_ab; 1.
DR PFAM; PF00306; ATP-synt_ab_C; 1.
DR Hydrolase; ATP synthesis; CF(1); ATP-binding;
KW Hydrogen ion transport.
FT NP_BIND 170 177 ATP (POTENTIAL).
SQ SEQUENCE 485 AA; 53034 MW; 07216783 CRC32;

Query Match 61.9%; Score 39; DB 1; Length 485;
Best Local Similarity 57.1%; Pred. No. 1.5;
Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 YRLAIRIXRILLRY 14
||:| : ||| ||
Db 376 YRVAQEVIRILQRY 389

RESULT 3
ATPB_MYCTU ID ATPB_MYCTU STANDARD; PRT; 486 AA.
AC Q10593;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE ATP SYNTHASE BETA CHAIN [EC 3.6.1.34].
GN ATPD OR RV1310 OR MTCY373.30.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-H37RV;
RX MEDLINE; 98295987.
RA COLE S.T., BROCH R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
RA GORDON S.V., EIGLMER K., GAS S., BARRY C.E. III, TEKAIA F.,
RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
RA DAVIES R., DEVLIN K., FELTWEILL T., GENTLES S., HAMLIN N., HOLROYD S.,
RA HORNSBY T., JAGELS K., KROGH A., MCLEAN J.J., MOULE S., MURPHY L.,
RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAM M.A., ROGERS J.,
RA RUTTER S., SEEGER K., SKELTON S., SQUARES S., SOARES R., SULSTON J.E.,
RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
RA "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence."
RL Nature 393:537-544(1998).
CC -!- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON
CC GRADIENT ACROSS THE MEMBRANE. THE BETA CHAIN IS THE CATALYTIC
CC SUBUNIT.
CC -!- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC
CC CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
CC HAS THREE MAIN SUBUNITS: A, B AND C. (BY SIMILARITY).
CC -!- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.

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CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; Z73419; CAA97743.1; -
DR HSSP; P07677; 1SKY.
DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
DR PFAM; PF00006; ATP-synt_ab; 1.
DR PFAM; PF00306; ATP-synt_ab_C; 1.
DR Hydrolase; ATP synthesis; CF(1); ATP-binding;
KW Hydrogen ion transport.
FT NP_BIND 171 178 ATP (POTENTIAL).
SQ SEQUENCE 486 AA; 53094 MW; A8001B2F CRC32;

Query Match 61.9%; Score 39; DB 1; Length 486;
Best Local Similarity 57.1%; Pred. No. 1.5;
Matches 8; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

Qy 1 YRLAIRIXRILLRY 14
||:| : ||| ||
Db 377 YRVAQEVIRILQRY 390

RESULT 4
UREF_HAEIN ID UREF_HAEIN STANDARD; PRT; 194 AA.
AC P44395;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE UREASE ACCESSORY PROTEIN UREF.
GN UREF OR HI0537.
OS Haemophilus influenzae.
OC Bacteria; Proteobacteria; gamma subdivision; Pasteurellaceae;
OC Haemophilus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-RD / KW20;
RX MEDLINE; 95350630.
RA FLEISCHMANN R.D., ADAMS M.D., WHITE O., CLAYTON R.A., KIRKNESS E.F.,
RA KERLAVAGE A.R., BULT C.J., TOMB J.-F., DOUGHERTY B.A., MERRICK J.M.,
RA MCKENNEY K., SUTTON G., FITZHUGH W., FIELDS C.A., GOCAYNE J.D.,
RA SCOTT J.D., SHIRLEY R., LIU L.-I., GLODEK A., KELLEY J.M.,
RA WEIDMAN J.F., PHILLIPS C.A., SPRIGGS T., HEDBLOM E., COTTON M.D.,
RA UTTERBACK T.R., HANNA M.C., NGUYEN D.T., SAUDEK D.M., BRANDON R.C.,
RA FINE L.D., FRITCHMAN J.L., FUHRMANN J.L., GEOGHAGEN N.S.M.,
RA GNEHM C.L., McDONALD L.A., SMALL K.V., FRASER C.M., SMITH H.O.,
RA VENTER J.C.;
RA "Whole-genome random sequencing and assembly of Haemophilus
RT influenzae RD."
RL Science 269:496-512(1995).
CC -!- FUNCTION: PROBABLY FACILITATING NICKEL INCORPORATION.
CC -!- SIMILARITY: BELONGS TO THE UREF FAMILY.
CC -----
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CC -----
DR EMBL; U32736; AAC22195.1; -
DR TIGR; HI0537; -
KW Nickel.
SQ SEQUENCE 194 AA; 21960 MW; 44EB0D8E CRC32;

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Query Match 60.3%; Score 38; DB 1; Length 194;
Best Local Similarity 35.7%; Pred. No. 0.9;
Matches 5; Conservative 6; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILRY 14
DB 64 FXLGVRLKIFIRY 77

RESULT 5
ATPB_HERAU STANDARD; PRT; 471 AA.
AC P42466;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE ATP SYNTHASE BETA CHAIN (EC 3.6.1.34).
GN ATPD.
OS Herpetosiphon aurantiacus (Herpetosiphon giganteus).
OC Bacteria; Green non-sulfur bacteria; Chloroflexaceae group;
OC Herpetosiphon.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=HPGAL;
RX MEDLINE; 94368062.
RA KLUGBAUER S., NEUMAYER J., KLUGBAUER N., BROCKMANN E., ROLLER C.,
RA KLUGBAUER S., REPTZ K., SCHACHTNER I., LUDVIGSEN A.,
RA BACHLEITNER M., FISCHER U., SCHLEIFER K.H.;
RT "Phylogenetic relationships of Bacteria based on comparative sequence
RT analysis of elongation factor Tu and ATP-synthase beta-subunit
RT genes."
RL Antonie van Leeuwenhoek 64:285-305(1993).
CC -!- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON
CC GRADIENT ACROSS THE MEMBRANE. THE BETA CHAIN IS THE CATALYTIC
CC SUBUNIT.
CC -!- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC
CC CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
CC HAS THREE MAIN SUBUNITS: A, B AND C.
CC -!- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
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CC -----
DR EMBL; X76876; CAA54203.1; -
DR HSSP; P07677; 1SKY.
DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
DR PFAM; PF00006; ATP-synt_ab; 1.
DR PFAM; PF00306; ATP-synt_ab_C; 1.
KW Hydrolyase; ATP synthesis; CF(1); ATP-binding;
KW Hydrogen ion transport.
FT NP_BIND 152 159 ATP (BY SIMILARITY).
SQ SEQUENCE 471 AA; 51057 MW; AAE39561 CRC32;

Query Match 60.3%; Score 38; DB 1; Length 471;
Best Local Similarity 50.0%; Pred. No. 2.3;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILRY 14
DB 364 YRVATEVQRMQLRY 377

RESULT 6
ATPB_AQUAE STANDARD; PRT; 478 AA.
AC P42466;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE ATP SYNTHASE BETA CHAIN (EC 3.6.1.34).
GN ATPD.
OS Aquifex pyrophilus.
OC Bacteria; Aquificales; Aquificaceae; Aquifex.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=DSM 6858;
RX MEDLINE; 98248216.
RA LUDWIG W., STRUNK O., KLUGBAUER S., KLUGBAUER N., WEIZENEGGER M.,
RA NEUMAYER J., BACHLEITNER M., SCHLEIFER K.H.;
RT "Bacterial phylogeny based on comparative sequence analysis."
RT Electrophoresis 19:554-568(1998).
RL
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CC -!- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON
 CC GRADIENT ACROSS THE MEMBRANE. THE BETA CHAIN IS THE CATALYTIC
 CC SUBUNIT.
 CC -!- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS. CF(1) - THE CATALYTIC
 CC CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
 CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
 CC HAS THREE MAIN SUBUNITS: A, B AND C.
 CC -!- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
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 CC
 CC EMBL: Y15786; CAA75780.1; -
 CC PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
 CC PFAM; PF00006; ATP-synt_ab; 1.
 CC PFAM; PF00306; ATP-synt_ab_C; 1.
 CC Hydrolase; ATP synthesis; CF(1); ATP-binding;
 CC Hydrogen ion transport.
 CC NP_BIND 163 170 ATP (POTENTIAL).
 CC SEQUENCE 478 AA: 53393 MW: 8f8adb9 CRC32:
 CC
 CC Query Match 60.3%; Score 38; DB 1; Length 478;
 CC Best Local Similarity 50.0%; Pred. No. 2.4; Indels 0; Gaps 0;
 CC Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;
 CC
 CC QY 1 YRLAIRXIRLLRY 14
 CC I : : : : :
 CC DB 369 YEYAVEVKKIRLQRY 382
 CC
 CC RESULT 8
 CC CA04_CANMA STANDARD; PRT; 708 AA.
 CC ID CA04_CANMA
 CC AC P05335;
 CC DT 01-NOV-1988 (Rel. 09, Created)
 CC DT 01-MAY-1992 (Rel. 22, Last sequence update)
 CC DT 01-MAY-1992 (Rel. 22, Last annotation update)
 CC DE ACYL-COENZYME A OXIDASE POX4 (EC 1.3.3.6) (ACYL-COA OXIDASE) (AOX).
 CC GN POX4.
 CC OS Candida maltosa (Yeast).
 CC OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 CC OC Candidaceae; Candida.
 CC [1]
 CC RN SEQUENCE FROM N.A.
 CC RP STRAIN=ATCC 20184;
 CC RX MEDLINE: 88124223.
 CC RA HILL D.E., BOULAY R., ROGERS D.;
 CC RT "Complete nucleotide sequence of the peroxisomal acyl CoA oxidase
 CC from the alkane-utilizing yeast Candida maltosa.";
 CC RL Nucleic Acids Res. 16:365-366(1988).
 CC -!- CATALYTIC ACTIVITY: ACYL-COA + O(2) = TRANS-2,3-DEHYDROACYL-COA +
 CC H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
 CC FROM 8 TO 18).
 CC -!- COFACTOR: FAD.
 CC -!- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
 CC SYSTEM.
 CC -!- SUBUNIT: HOMOOCTAMER.
 CC -!- SUBCELLULAR LOCATION: PEROXISOMAL.
 CC DR PIR; A28584; OXCRAX.
 CC KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
 CC Multigene family.
 CC NON_TER 1
 CC SEQUENCE 502 AA: 55528 MW: EACE80C4 CRC32:
 CC
 CC Query Match 57.1%; Score 36; DB 1; Length 502;
 CC Best Local Similarity 50.0%; Pred. No. 6.4; Indels 0; Gaps 0;
 CC Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;
 CC
 CC QY 1 YRLAIRXIRLLRY 14
 CC I : : : : :
 CC DB 101 YRLMRSTIALRY 114
 CC
 CC RESULT 10
 CC CA02_CANTR STANDARD; PRT; 708 AA.
 CC ID CA02_CANTR
 CC AC P06598;
 CC DT 01-JAN-1988 (Rel. 06, Created)
 CC DT 01-NOV-1988 (Rel. 09, Last sequence update)
 CC DT 01-NOV-1997 (Rel. 35, Last annotation update)
 CC DE ACYL-COENZYME A OXIDASE II (EC 1.3.3.6) (ACYL-COA OXIDASE) (PXP-4).
 CC GN AOX OR POX-4.
 CC OS Candida tropicalis (Yeast).
 CC OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 CC OC Candidaceae; Candida.

DR EMBL; X06721; CAA29901.1; -
 DR PIR; A29441; OXCKPM.
 KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
 KW Multigene family.
 FT INIT_MET 0 BY SIMILARITY.
 SQ SEQUENCE 708 AA: 78242 MW: D5E344D2 CRC32:
 CC
 CC Query Match 60.3%; Score 38; DB 1; Length 708;
 CC Best Local Similarity 50.0%; Pred. No. 3.6; Indels 0; Gaps 0;
 CC Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;
 CC
 CC QY 1 YRLAIRXIRLLRY 14
 CC I : : : : :
 CC DB 307 YRLMRSTIALRY 320
 CC
 CC RESULT 9
 CC CA03_CANTR STANDARD; PRT; 502 AA.
 CC ID CA03_CANTR
 CC AC P11355;
 CC DT 01-JUL-1989 (Rel. 11, Created)
 CC DT 01-JUL-1989 (Rel. 11, Last sequence update)
 CC DT 01-NOV-1997 (Rel. 35, Last annotation update)
 CC DE ACYL-COENZYME A OXIDASE POX4-2 (EC 1.3.3.6) (ACYL-COA OXIDASE)
 CC (FRAGMENT).
 CC GN POX4-2.
 CC OS Candida tropicalis (Yeast).
 CC OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 CC OC Candidaceae; Candida.
 CC [1]
 CC RN SEQUENCE FROM N.A.
 CC RP MEDLINE: 87280361.
 CC RA SMALL G.M., LAZAROW P.B.;
 CC RT "Import of the carboxy-terminal portion of acyl-CoA oxidase into
 CC peroxisomes of Candida tropicalis.";
 CC RL J. Cell Biol. 105:247-250(1987).
 CC -!- CATALYTIC ACTIVITY: ACYL-COA + O(2) = TRANS-2,3-DEHYDROACYL-COA +
 CC H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
 CC FROM 8 TO 18).
 CC -!- COFACTOR: FAD.
 CC -!- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
 CC SYSTEM.
 CC -!- SUBUNIT: HOMOOCTAMER.
 CC -!- SUBCELLULAR LOCATION: PEROXISOMAL.
 CC DR PIR; A28584; OXCRAX.
 CC KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
 CC Multigene family.
 CC NON_TER 1
 CC SEQUENCE 502 AA: 55528 MW: EACE80C4 CRC32:
 CC
 CC Query Match 57.1%; Score 36; DB 1; Length 502;
 CC Best Local Similarity 50.0%; Pred. No. 6.4; Indels 0; Gaps 0;
 CC Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;
 CC
 CC QY 1 YRLAIRXIRLLRY 14
 CC I : : : : :
 CC DB 101 YRLMRSTIALRY 114
 CC
 CC RESULT 10
 CC CA02_CANTR STANDARD; PRT; 708 AA.
 CC ID CA02_CANTR
 CC AC P06598;
 CC DT 01-JAN-1988 (Rel. 06, Created)
 CC DT 01-NOV-1988 (Rel. 09, Last sequence update)
 CC DT 01-NOV-1997 (Rel. 35, Last annotation update)
 CC DE ACYL-COENZYME A OXIDASE II (EC 1.3.3.6) (ACYL-COA OXIDASE) (PXP-4).
 CC GN AOX OR POX-4.
 CC OS Candida tropicalis (Yeast).
 CC OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 CC OC Candidaceae; Candida.

[1]
RN SEQUENCE FROM N.A.
RP STRAIN-ATCC 20336 / PK233;
RX MEDLINE: 87248070.
RA MURRAY W.W., RACHUBINSKI R.A.;
RT "The primary structure of a peroxisomal fatty acyl-CoA oxidase from
the yeast *Candida tropicalis* pk233.";
RL Gene 51:119-128(1987).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-ATCC 20336 / PK233;
RX MEDLINE: 86149279.
RA OKAZAKI K., TAKECHI T., KAMBARA N., FUKUYAMA I., KAMIRYO T.;
RT "Two acyl-coenzyme A oxidases in peroxisomes of the yeast *Candida*
tropicalis: primary structures deduced from genomic DNA sequence.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:1232-1236(1986).
RN [3]
RP SEQUENCE OF 208-709 FROM N.A.
RC STRAIN-RR1;
RX MEDLINE: 87280361.
RA SMALL G.M., LAZAROW P.B.;
RT "Import of the carboxy-terminal portion of acyl-CoA oxidase into
peroxisomes of *Candida tropicalis*.";
RL J. Cell Biol. 105:247-250(1987).
CC -!- CATALYTIC ACTIVITY: ACYL-COA + O(2) = TRANS-2,3-DEHYDROACYL-COA +
H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
FROM 8 TO 18).
CC -!- COFACTOR: FAD.
CC -!- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
SYSTEM.
CC -!- SUBUNIT: HOMOOCTAMER.
CC -!- SUBCELLULAR LOCATION: PEROXISOMAL.
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CC
CC EMBL: M16193; AAA34322.1; -
DR EMBL: M12160; AAA34362.1; -
DR EMBL: Y00623; CAA68660.1; -
DR EMBL: Y00623; CAA68661.1; ALT_INIT.
DR EMBL: Y00623; CAA68662.1; ALT_INIT.
DR PIR: A25123; OXCKX4.
DR PIR: A29047; OXCKX.
KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
KW Multigene family.
FT INIT_MET 0
SQ SEQUENCE 708 AA; 79041 MW; D97A4EC8 CRC32;

Query Match 57.1%; Score 36; DB 1; Length 708;
Best Local Similarity 50.0%; Pred. NO. 9.2;
Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 YRLAIRXIRILLRY 14
||: | | | |

DB 307 YRLAIRXIRILLRY 320

RESULT 11

CAO4_CANTR STANDARD; PRT: 723 AA.
ID CAO4_CANTR
AC P11356;
DT 01-JUL-1989 (Rel. 11, Created)
DT 01-MAY-1992 (Rel. 22, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE ACYL-COENZYME A OXIDASE PXP-2 (EC 1.3.3.6) (ACYL-COA OXIDASE).
GN POX2.
OS *Candida tropicalis* (Yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Candidaceae; *Candida*.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE: 88084444.
RA OKAZAKI K., TAN H., FUKUYAMA I., KUBOTA I., KAMIRYO T.;
RT "Peroxisomal acyl-coenzyme A oxidase multigene family of the yeast
Candida tropicalis; nucleotide sequence of a third gene and its
protein product.";
RL Gene 58:37-44(1987).
CC -!- CATALYTIC ACTIVITY: ACYL-COA + O(2) = TRANS-2,3-DEHYDROACYL-COA +
H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
FROM 8 TO 18).
CC -!- COFACTOR: FAD.
CC -!- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
SYSTEM.
CC -!- SUBUNIT: HOMOOCTAMER.
CC -!- SUBCELLULAR LOCATION: PEROXISOMAL.
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CC
CC EMBL: M18259; AAA34361.1; -
DR PIR: A27331; OXCRP2.
KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
KW Multigene family.
FT INIT_MET 0
SQ SEQUENCE 723 AA; 81804 MW; 60C2D2B7 CRC32;

Query Match 55.6%; Score 35; DB 1; Length 723;
Best Local Similarity 50.0%; Pred. No. 15;
Matches 7; Conservative 1; Mismatches 6; Indels 0; Gaps 0;

QY 1 YRLAIRXIRILLRY 14
||: | | | |

DB 329 YRLAIRXIRILLRY 342

RESULT 12

CAO2_CANMA STANDARD; PRT: 724 AA.
ID CAO2_CANMA
AC Q00468;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE ACYL-COENZYME A OXIDASE POX2 (EC 1.3.3.6) (ACYL-COA OXIDASE) (AOX).
GN POX2.

OS *Candida maltosa* (Yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Candidaceae; *Candida*.
RN [1]

RP SEQUENCE FROM N.A.
RX STRAIN-1AM 12247;
RX MEDLINE: 96144267.

RA MASUDA Y., PARK S.M., OHTA A., TAKAGI M.;
RT "Cloning and characterization of the POX2 gene in *Candida maltosa*.";
RL Gene 167:157-161(1995).

CC -!- CATALYTIC ACTIVITY: ACYL-COA + O(2) = TRANS-2,3-DEHYDROACYL-COA +
H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
FROM 8 TO 18).
CC -!- COFACTOR: FAD.
CC -!- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
SYSTEM.
CC -!- SUBUNIT: HOMOOCTAMER.
CC -!- SUBCELLULAR LOCATION: PEROXISOMAL.
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CC
CC EMBL: M18259; AAA34361.1; -
DR PIR: A27331; OXCRP2.
KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
KW Multigene family.
FT INIT_MET 0
SQ SEQUENCE 723 AA; 81804 MW; 60C2D2B7 CRC32;

```

CC H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
CC FROM 8 TO 18).
CC -|- COFACTOR: FAD.
CC -|- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
CC SYSTEM.
CC -|- SUBUNIT: HOMOOCTAMER (BY SIMILARITY).
CC -|- SUBCELLULAR LOCATION: PEROXISOMAL.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: D21228; BAA04761.1; -.
CC DR Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
CC KW Multigene family.
CC SQ SEQUENCE 724 AA; 82273 MW; 1AE92F21 CRC32;
CC -----
CC Query Match 55.68; Score 35; DB 1; Length 724;
CC Best Local Similarity 50.08; Pred. No. 15;
CC Matches 7; Conservative 1; Mismatches 6; Indels 0; Gaps 0;
CC -----
CC QY 1 YRLAIRIXRILLRY 14
CC II: | | | | |
CC DB 330 YRICARTTIALRY 343
CC -----
CC RESULT 13
CC ACCA_AQVAE
CC ID ACCA_AQVAE STANDARD; PRT; 323 AA.
CC AC 067260;
CC DT 15-JUL-1999 (Rel. 38, Created)
CC DT 15-JUL-1999 (Rel. 38, Last sequence update)
CC DT 15-DEC-1999 (Rel. 39, Last annotation update)
CC DE ACETYL-COENZYME A CARBOXYLASE CARBOXYL TRANSFERASE SUBUNIT ALPHA
CC (EC 6.4.1.2)
CC GN ACQA OR AQ_1206.
CC OS Aquifex aeolicus.
CC OC Bacteria; Aquificales; Aquificaceae; Aquifex.
CC RN [1]
CC RP SEQUENCE FROM N.A.
CC RC STRAIN-VF5.
CC RX MEDLINE; 98196666.
CC RA DECKERT G., WARREN P.V., GAASTERLAND T., YOUNG W.G., LENOX A.L.,
CC GRAHAM D.E., OVERBECK R., SNEAD M.A., KELLER M., AUJAY M., HUBER R.,
CC FELDMAN R.A., SHORT J.M., OLSON G.J., SWANSON R.V.;
CC RT "The complete genome of the hyperthermophilic bacterium Aquifex
CC aeolicus."
CC RL Nature 392:353-358(1998).
CC CC -|- FUNCTION: THIS PROTEIN IS A COMPONENT OF THE ACETYL COENZYME A
CC CARBOXYLASE COMPLEX; FIRST, BIOTIN CARBOXYLASE CATALYZES THE
CC CARBOXYLATION OF THE CARRIER PROTEIN AND THEN THE TRANSCARBOXYLASE
CC TRANSFERS THE CARBOXYL GROUP TO FORM MALONYL-COA (BY SIMILARITY).
CC -|- CATALYTIC ACTIVITY: CARBOXYBIOTIN CARBOXYL CARRIER PROTEIN +
CC ACETYL-COA -> BIOTIN CARBOXYL CARRIER PROTEIN + MALONYL-COA.
CC -|- PATHWAY: FIRST STEP IN LONG-CHAIN FATTY ACID SYNTHESIS.
CC CC -|- SUBUNIT: ACETYL-COA CARBOXYLASE IS AN HETERODIMER OF BIOTIN
CC CARBOXYL CARRIER PROTEIN, BIOTIN CARBOXYLASE AND THE TWO SUBUNITS
CC OF CARBOXYL TRANSFERASE IN A 2:2 COMPLEX (BY SIMILARITY).
CC -|- SIMILARITY: TO THE C-TERMINUS OF MAMMALIAN PROPIONYL-COA
CC CARBOXYLASE BETA CHAIN.
CC -----
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CC -----
CC EMBL: AF000728; AAC07216.1; -.
CC DR Fatty acid biosynthesis; Ligase.
CC KW SEQUENCE 323 AA; 36198 MW; 4A96C81E CRC32;
CC SQ -----
CC Query Match 54.08; Score 34; DB 1; Length 323;
CC Best Local Similarity 42.98; Pred. No. 10;
CC Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;
CC -----
CC QY 1 YRLAIRIXRILLRY 14
CC II: | | | | |
CC DB 140 YRKAIKFKLAERY 153
CC -----
CC RESULT 14
CC NY5R_CANFA
CC ID NY5R_CANFA STANDARD; PRT; 446 AA.
CC AC 062729;
CC DT 15-DEC-1998 (Rel. 37, Created)
CC DT 15-DEC-1998 (Rel. 37, Last sequence update)
CC DT 15-DEC-1999 (Rel. 39, Last annotation update)
CC DE NEUROPEPTIDE Y RECEPTOR TYPE 5 (NPY5-R) (NPY-Y5 RECEPTOR) (Y5
CC RECEPTOR) (NPY5).
CC GN NPY5R OR NPY5.
CC OS Canis familiaris (Dog).
CC OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
CC Eutheria; Carnivora; Fissipedia; Canidae; Canis.
CC RN [1]
CC RP SEQUENCE FROM N.A.
CC RX MEDLINE; 99017379.
CC RA BOROWSKY B., WALKER M.W., BARD J., WEINSHANK R.L., LAZ T.M.,
CC VAYASSE P., BRANCHER T.A., GERALD C.;
CC RT "Molecular biology and pharmacology of multiple NPY Y5 receptor
CC species homologs."
CC RL Regul. Pept. 75:45-53(1998).
CC CC -|- FUNCTION: RECEPTOR FOR NEUROPEPTIDE Y AND PEPTIDE YY. THE ACTIVITY
CC OF THIS RECEPTOR IS MEDIATED BY G PROTEINS THAT INHIBITS ADENYLATE
CC CYCLASE ACTIVITY. SEEMS TO BE ASSOCIATED WITH FOOD INTAKE. COULD
CC BE INVOLVED IN FEEDING DISORDERS (BY SIMILARITY).
CC -|- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
CC -|- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
CC -|- HIGHEST TO TACHIKININS RECEPTORS.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: AF049328; AAC17838.1; -.
CC DR PROSITE; PS00237; G-PROTEIN_RECEPTOR; FALSE_NEG.
CC DR PFAM; PF00001; 7tm1; 1.
CC KW G-protein coupled receptor; Transmembrane; Glycoprotein;
CC Phosphorylation; Lipoprotein; Palmitate.
CC FT DOMAIN 1 39 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 41 62 1 (POTENTIAL).
CC FT DOMAIN 63 74 CYTOPLASMIC (POTENTIAL).
CC FT TRANSMEM 75 95 2 (POTENTIAL).
CC FT DOMAIN 96 115 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 116 137 3 (POTENTIAL).
CC FT DOMAIN 138 157 CYTOPLASMIC (POTENTIAL).
CC FT TRANSMEM 158 178 4 (POTENTIAL).
CC FT DOMAIN 179 210 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 211 232 5 (POTENTIAL).
CC FT DOMAIN 233 311 CYTOPLASMIC (POTENTIAL).
CC FT TRANSMEM 369 391 6 (POTENTIAL).
CC FT DOMAIN 392 404 EXTRACELLULAR (POTENTIAL).
CC FT TRANSMEM 405 428 7 (POTENTIAL).
CC FT DOMAIN 429 446 CYTOPLASMIC (POTENTIAL).
CC FT CARBOHYD 10 10 POTENTIAL.

```

FT CARBOHYD 17 17 POTENTIAL.
 FT DISULFID 114 198 BY SIMILARITY.
 FT LIPID 442 442 PALMITATE (POTENTIAL).
 SQ SEQUENCE 446 AA; 51012 MW; 7D2CD74A CRC32;

Query Match 54.0%; Score 34; DB 1; Length 446;
 Best Local Similarity 42.9%; Pred. NO. 14;
 Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Oy 1 YRLAIRIXRILLRY 14
 ||:| | :|:|
 Db 207 YRIAFITSLLLVQY 220

RESULT 15

NY5R_PIG STANDARD; PRT; 446 AA.
 AC O97969;
 DT 15-DEC-1999 (Rel. 39, Created)
 DT 15-DEC-1999 (Rel. 39, Last sequence update)
 DT 15-DEC-1999 (Rel. 39, Last annotation update)
 DE NEUROPEPTIDE Y RECEPTOR TYPE 5 (NPY5-R) (NPY-Y5 RECEPTOR) (Y5 RECEPTOR).
 GN NPY5R OR NPYR5.
 OS Sus scrofa (Pig).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA WRAITH A., TORNSTEN A., CHARDON P., HARBITZ I., CHOWDHARY B.P.,
 RA ANDERSSON L., LARHAMMAR D.;
 RT "Porcine NPY receptors NPY1R, NPY2R and NPY5R: cloning, mapping and
 RT comparative analysis."
 RL Submitted (NOV-1998) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-LWD; TISSUE=KIDNEY;
 RA ITO Y., MINEZAWA M.;
 RT "Sus scrofa gene for neuropeptide Y receptor type 5, complete cds."
 RL Submitted (OCT-1998) to the EMBL/GenBank/DBJ databases

CC -!- FUNCTION: RECEPTOR FOR NEUROPEPTIDE Y AND PEPTIDE YY. THE ACTIVITY
 CC OF THIS RECEPTOR IS MEDIATED BY G PROTEINS THAT INHIBITS ADENYLATE
 CC CYCLASE ACTIVITY. SEEMS TO BE ASSOCIATED WITH FOOD INTAKE. COULD
 CC BE INVOLVED IN FEEDING DISORDERS (BY SIMILARITY).
 CC -!- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN.
 CC -!- SIMILARITY: BELONGS TO FAMILY 1 OF G-PROTEIN COUPLED RECEPTORS.
 CC HIGHEST TO TACHYKININS RECEPTORS.
 CC -----
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 CC -----

DR EMBL; AF106083; AAD13778.1; -
 DR EMBL; AB019185; BAA34055.1; -
 DR PROSITE; PS00237; G_PROTEIN_RECEPTOR; FALSE_NEG.
 KW G-protein coupled receptor; Transmembrane; Glycoprotein;
 KW Phosphorylation; Lipoprotein; Palmitate.
 FT DOMAIN 1 39 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 41 62 1 (POTENTIAL).
 FT DOMAIN 63 74 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 75 95 2 (POTENTIAL).
 FT DOMAIN 96 115 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 116 137 3 (POTENTIAL).
 FT DOMAIN 138 157 CYTOPLASMIC (POTENTIAL).
 FT TRANSMEM 158 178 4 (POTENTIAL).
 FT DOMAIN 179 210 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 211 232 5 (POTENTIAL).
 FT DOMAIN 233 311 CYTOPLASMIC (POTENTIAL).

FT TRANSMEM 369 391 6 (POTENTIAL).
 FT DOMAIN 392 404 EXTRACELLULAR (POTENTIAL).
 FT TRANSMEM 405 428 7 (POTENTIAL).
 FT DOMAIN 429 446 CYTOPLASMIC (POTENTIAL).
 FT CARBOHYD 10 10 POTENTIAL.
 FT CARBOHYD 17 17 POTENTIAL.
 FT DISULFID 114 198 BY SIMILARITY.
 FT LIPID 442 442 PALMITATE (POTENTIAL).
 SQ SEQUENCE 446 AA; 50474 MW; 79A4E2F3 CRC32;

Query Match 54.0%; Score 34; DB 1; Length 446;
 Best Local Similarity 42.9%; Pred. NO. 14;
 Matches 6; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

Oy 1 YRLAIRIXRILLRY 14
 ||:| | :|:|
 Db 207 YRIAFITSLLLVQY 220

Search completed: February 8, 2000, 00:59:58
 Job time: 3787 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:48 : Search time 209.03 Seconds
(without alignments)
4.644 Million cell updates/sec

Title: US-08-653-294-20

Perfect score: 63

Sequence: 1 YRLAIRIXRILLRY 14

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 225878 segs, 69334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database :

SPTREMBL_12.*

1: sp_archaea.*

2: sp_bacteria.*

3: sp_fungi.*

4: sp_human.*

5: sp_invertebrate.*

6: sp_mammal.*

7: sp_mmc.*

8: sp_organelle.*

9: sp_phage.*

10: sp_plant.*

11: sp_rodent.*

12: sp_virus.*

13: sp_vertebrate.*

14: sp_unclassified.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	38	60.3	133	2 Q92903	Q92903 chlamydia p
2	38	60.3	2810	5 Q20456	Q20456 caenorhabdi
3	37	58.7	1592	5 Q45019	Q45019 caenorhabdi
4	36	57.1	96	1 Q29637	Q29637 archaeoglob
5	36	57.1	248	3 Q74720	Q74720 debaryomyce
6	36	57.1	353	2 Q70073	Q70073 agrobacteri
7	36	57.1	3131	3 Q00869	Q00869 fusarium sc
8	35	55.6	265	2 Q44541	Q44541 azotobacter
9	35	55.6	402	2 Q57090	Q57090 corynebacte
10	35	55.6	404	10 Q9X1L2	Q9X1L2 arabidopsis
11	35	55.6	469	2 Q50159	Q50159 streptococc
12	34	54.0	44	10 Q49025	Q49025 gracilaria
13	34	54.0	157	1 Q9YC92	Q9YC92 aeropyrum p
14	34	54.0	170	2 P94457	P94457 bacillus st.
15	34	54.0	276	1 Q9YFD0	Q9YFD0 aeropyrum p
16	34	54.0	372	13 Q93237	Q93237 cyprinus ca
17	34	54.0	468	2 Q9ZJ01	Q9ZJ01 streptococc
18	34	54.0	1021	4 Q15451	Q15451 homo sapien
19	34	54.0	1251	4 Q15450	Q15450 homo sapien
20	33	52.4	149	2 Q9X1G5	Q9X1G5 thermotoga

21	33	52.4	485	2	P71815	P71815 mycobacteri
22	33	52.4	499	10	O04376	O04376 arabidopsis
23	33	52.4	532	2	P72007	P72007 mycobacteri
24	33	52.4	719	3	Q9Y7B1	Q9Y7B1 pichia past
25	33	52.4	2630	12	O55319	O55319 acyrthosiph
26	33	52.4	3074	12	O08534	O08534 sugar beet
27	32.5	51.6	42	11	O54741	O54741 mus musculu
28	32	50.8	339	1	Q9YA89	Q9YA89 aeropyrum p
29	32	50.8	422	12	O88530	O88530 turkey herp
30	32	50.8	591	2	Q9Z8G2	Q9Z8G2 chlamydia p
31	32	50.8	785	10	O23072	O23072 arabidopsis
32	32	50.8	2962	5	Q93326	Q93326 caenorhabdi
33	32	50.8	3070	12	O89906	O89906 beet yellow
34	31	49.2	76	3	Q06739	Q06739 saccharomyc
35	31	49.2	169	11	O70180	O70180 rattus norv
36	31	49.2	185	2	O55192	O55192 synecocyst
37	31	49.2	192	3	O13610	O13610 schizosacch
38	31	49.2	211	5	Q9XU58	Q9XU58 caenorhabdi
39	31	49.2	220	1	Q9Y961	Q9Y961 aeropyrum p
40	31	49.2	223	2	Q9ZIN6	Q9ZIN6 staphylococ
41	31	49.2	224	2	O83422	O83422 treponema p
42	31	49.2	275	5	Q93780	Q93780 caenorhabdi
43	31	49.2	277	2	O52027	O52027 pseudomonas
44	31	49.2	294	8	Q02676	Q02676 podospora a
45	31	49.2	331	1	O58520	O58520 pyrococcus

ALIGNMENTS

RESULT 1

Q92903 ID Q92903 PRELIMINARY; PRT; 133 AA.
AC Q92903:
DT 01-MAY-1999 (TREMBlrel. 10, Created)
DT 01-MAY-1999 (TREMBlrel. 10, Last sequence update)
DE 01-MAY-1999 (TREMBlrel. 10, Last annotation update)
DE HYPOTHETICAL 16.1 KD PROTEIN.
GN CPN0181.
OS Chlamydia pneumoniae.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydiaophila.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=CWLO29;
RA KALMAN S., MITCHELL W., MARATHE R., LAMMEL C., FAN J., OLINGER L.,
RA GRIMWOOD J., DAVIS R.W., STEPHENS R.S.;
RT "Comparative Genomes of chlamydia pneumoniae and C. trachomatis.";
RL Submitted (DEC-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL; AE001604; AAD18334.1; -
KW Hypothetical protein.
SQ SEQUENCE 133 AA; 16132 MW; 8DA54C6A CRC32;

Query Match 60.3%; Score 38; DB 2; Length 133;
Best Local Similarity 61.5%; Pred. No. 4.3;
Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 RLAIKXIRILLRY 14

Db 88 RIPWRLKIRILLRY 100

RESULT 2

Q20456 ID Q20456 PRELIMINARY; PRT; 2810 AA.
AC Q20456:
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1998 (TREMBlrel. 08, Last sequence update)
DE 01-MAY-1999 (TREMBlrel. 10, Last annotation update)
DE HUM-4 PROTEIN.

GN HUM-4.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;

OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA COTTAGE A.;
 RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 94150718.
 RA WILSON R., AINSKOUGH R., ANDERSON K., BAYNES C., BERKS M.,
 RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
 RA CRAYTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
 RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,
 RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
 RA LIGHTNING J., LLOYD C., McMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
 RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
 RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
 RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
 RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 RT elegans.";
 RL Nature 368:32-38(1994).
 DR EMBL; 265563; CAA91469.1; -;
 DR PFAM; PF00612; IQ_2; myosin_head; 4.
 DR PFAM; PF00063; myosin_head; 4.
 DR PFAM; PF00784; MYTH4; 2.
 SQ SEQUENCE 2810 AA; 323526 MW; 6274286C CRC32;

Query Match 60.3%; Score 38; DB 5; Length 2810;
 Best Local Similarity 35.7%; Pred. NO. 86;
 Matches 5; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLRY 14
 :||:| :||:|
 Db 1102 FRLSVEFKILAY 1115

RESULT 3
 ID 045019 PRELIMINARY; PRT; 1592 AA.
 AC 045019;
 DT 01-JUN-1998 (TRENBLrel. 06, Created)
 DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE ZC123.3 PROTEIN.
 GN ZC123.3.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditidae;
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BRISTOL N2;
 RX MEDLINE; 99069613.
 RA NONE;
 RT "Genome sequence of the nematode C. elegans: a platform for
 RT investigating biology. The C. elegans Sequencing Consortium.";
 RT Science 282:2012-2018(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BRISTOL N2;
 RA BRADSHAW H., GRAVES T., BIEWALD T.;
 RT "The sequence of C. elegans cosmid ZC123.";
 RL Submitted (JAN-1998) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=BRISTOL N2;
 RA WATERSTON R.;
 RL Submitted (MAY-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- SUBCELLULAR LOCATION: NUCLEAR (BY SIMILARITY).
 DR EMBL; AF043706; AAB97603.2; -;
 DR PROSITE; PS00027; HOMEBOX.1; 1.
 DR PROSITE; PS00028; ZINC_FINGER_C2H2; 6.
 DR PFAM; PF00096; zf-C2H2; 2.

KW Homeobox; DNA-binding; Nuclear protein; Zinc-finger; Metal-binding.
 SQ SEQUENCE 1592 AA; 178053 MW; BAFDE8CE CRC32;

Query Match 58.7%; Score 37; DB 5; Length 1592;
 Best Local Similarity 72.7%; Pred. NO. 77;
 Matches 8; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLAIKXIRILL 12
 :||:| :||:|
 Db 477 RYAIRLIRILL 487

RESULT 4
 ID 029637 PRELIMINARY; PRT; 96 AA.
 AC 029637;
 DT 01-JAN-1998 (TRENBLrel. 05, Created)
 DT 01-JAN-1998 (TRENBLrel. 05, Last sequence update)
 DT 01-AUG-1998 (TRENBLrel. 07, Last annotation update)
 DE CONSERVED HYPOTHETICAL PROTEIN.
 GN AF0618.
 OS Archaeoglobus fulgidus.
 OC Archaea; Euryarchaeota; Archaeoglobales; Archaeoglobaceae;
 OC Archaeoglobus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=VC-16 / DSM 4304 / ATCC 49558;
 RX MEDLINE; 98049343.
 RA KLENK H.-P., CLAYTON R.A., TOMB J.-F., WHITE O., NELSON K.E.,
 RA KECHUM K.A., DODSON R.J., GWINN M., HICKEY E.K., PETERSON J.D.,
 RA RICHARDSON D.L., KERLAVAGE A.R., GRAHAM D.E., KYRPIDES N.C.,
 RA FLEISCHMANN R.D., QUACKENBUSH J., LEE N.H., SUTTON G.G., GILL S.,
 RA KIRKNESS E.F., DOUGHERTY B.A., MCKENNEY K., ADAMS M.D., LOFTUS B.,
 RA PETERSON S., REICH C.I., MCNEIL L.K., BADGER J.H., GLODEK A., ZHOU L.,
 RA OVERBECK R., GOCAYNE J.D., WEIDMAN J.F., McDONALD L., UTTERBACK T.,
 RA COTTON M.D., SPRIGGS T., ARTIACH P., KATNE B.P., SYKES S.W.,
 RA SADON P.W., D'ANDREA K.P., BOWMAN C., FUJII C., GARLAND S.A.,
 RA MASON T.M., OLSEN G.J., FRASER C.M., SMITH H.O., WOESE C.R.,
 RA VENTER J.C.;
 RT "The complete genome sequence of the hyperthermophilic, sulphate-
 RT reducing archaeon Archaeoglobus fulgidus.";
 RL Nature 390:364-370(1997).
 DR EMBL; AE001062; AAB90622.1; -;
 DR TIGR; AF0618; -;
 KW Hypothetical protein.
 SQ SEQUENCE 96 AA; 10892 MW; 76C3565A CRC32;

Query Match 57.1%; Score 36; DB 1; Length 96;
 Best Local Similarity 61.5%; Pred. NO. 7.6;
 Matches 8; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILLR 13
 :||:| :||:|
 Db 51 YRLAIKISTELLK 63

RESULT 5
 ID 074720 PRELIMINARY; PRT; 248 AA.
 AC 074720;
 DT 01-NOV-1998 (TRENBLrel. 08, Created)
 DT 01-NOV-1998 (TRENBLrel. 08, Last sequence update)
 DT 01-NOV-1998 (TRENBLrel. 08, Last annotation update)
 DE LINEAR PLASMID PDH1, ORF1 AND ORF2, PARTIAL (FRAGMENT).
 OS Debaryomyces hansenii (Yeast) (Torulaspora hansenii).
 OG Plasmid pDH1.
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Debaryomyces.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC90624, CBS7848;

RA GUNGE N.;
 RL Submitted (OCT-1998) to the EMBL/GenBank/DBJ databases.
 [2]

RN SEQUENCE FROM N.A.
 RP STRAIN=ATCC90624, CBS7848;
 RX MEDLINE: 97344364.
 RA FUKUDA K., MAESUCHI M., TAKATA H., GUNGE N.;
 RT "The linear plasmid pDHL1 from Debaryomyces hansenii encodes a protein
 highly homologous to the pgkII-plasmid DNA polymerase.";
 RL yeast 13:613-620(1997).
 DR EMBL: AJ011124; CAA09498.1; -.
 KW Plasmid.
 FT NON_TER 248 248
 SQ SEQUENCE 248 AA; 27239 MW; 54CD45A4 CRC32;

Query Match 57.1%; Score 36; DB 3; Length 248;
 Best Local Similarity 75.0%; Pred. No. 19;
 Matches 9; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 YRLAIRXRILL 12
 |||||
 DB 4 YRLAIFASILL 15

RESULT 6
 ID 070073 PRELIMINARY; PRT; 353 AA.
 AC 070073;
 DT 01-AUG-1998 (TEMBLrel. 07, Created)
 DT 01-AUG-1998 (TEMBLrel. 07, Last sequence update)
 DT 01-AUG-1998 (TEMBLrel. 07, Last annotation update)
 DE PUTATIVE PERIPLASMIC PROTEIN CHTH.
 GN CHTH.
 OS Agrobacterium tumefaciens, and Agrobacterium sp.
 OG Plasmid pTichry5.
 OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
 OC Rhizobiaceae; Agrobacterium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=CHRY5;
 RA OGER P.M., VAUDEQUIN V., DESSAUX Y.;
 RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
 [2]
 RN SEQUENCE FROM N.A.
 RC STRAIN=ANT4;
 RA OGER P.M., VAUDEQUIN-DRANSART V., DESSAUX Y.;
 RL Submitted (APR-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF063242; AAC17432.1; -.
 DR EMBL: U67851; AAC12800.1; -.
 KW Plasmid.
 SQ SEQUENCE 353 AA; 38687 MW; 4B019D49 CRC32;

Query Match 57.1%; Score 36; DB 2; Length 353;
 Best Local Similarity 46.2%; Pred. No. 27;
 Matches 6; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 2 RLAIRXRILL 14
 :||:|:|
 DB 247 KLGLRGRVRLY 259

RESULT 7
 ID Q00869 PRELIMINARY; PRT; 3131 AA.
 AC Q00869;
 DT 01-NOV-1996 (TEMBLrel. 01, Created)
 DT 01-NOV-1996 (TEMBLrel. 01, Last sequence update)
 DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
 DE ENNTATIN SYNTHASE.
 GN ESNL.
 OS Fusarium scirpi.

OC Eukaryota; Fungi; Ascomycota; Euascomycetes; Pyrenomycetes;
 OC Hypocreales; Hypocreaceae; anamorphic Hypocreaceae; Fusarium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=LAMBOTTE ET FAUTREV;
 RA HAENSE A., SCHUBERT M., HERRMANN M., ZOCHER R.;
 RL Mol. Microbiol. 0:0-0(1992).
 DR EMBL: Z18755; CAA79245.1; -.
 DR MENDEL; 20784; Fuser; 3105; 20784.
 DR PROSITE; PS00455; AMP_BINDING; 2.
 DR PFAM; PF00501; AMP-binding; 2.
 DR PFAM; PF00668; DUF4; 2.
 DR PFAM; PF00550; PP-binding; 3.
 DR PRINTS; PR00154; AMPBINDING.
 SQ SEQUENCE 3131 AA; 346891 MW; 00949DB9 CRC32;

Query Match 57.1%; Score 36; DB 3; Length 3131;
 Best Local Similarity 53.8%; Pred. No. 2.3e+02;
 Matches 7; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLAIRXRILL 14
 :||:|:|
 DB 2637 KLAIRGRRLRH 2649

RESULT 8
 ID Q44541 PRELIMINARY; PRT; 265 AA.
 AC Q44541;
 DT 01-NOV-1996 (TEMBLrel. 01, Created)
 DT 01-NOV-1996 (TEMBLrel. 01, Last sequence update)
 DT 01-NOV-1999 (TEMBLrel. 12, Last annotation update)
 DE ORF 7.
 OS Azotobacter vinelandii.
 OC Bacteria; Proteobacteria; gamma subdivision; Azotobacteraceae;
 OC Azotobacter.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE: 89123097.
 RA JACOBSON M.R., BRIGLE K.E., BENNETT L.T., SETTERQUIST R.A.,
 RA WILSON M.S., CASH V.L., BEYNON J., NEWTON W.E., DEAN D.R.;
 RT "Physical and genetic map of the major nif gene cluster from
 Azotobacter vinelandii.";
 RL J. Bacteriol. 171:1017-1027(1989).
 DR EMBL: M20568; AAA64728.1; -.
 DR PROSITE; PS00101; HEXAPEP_TRANSFERASES; 1.
 DR PFAM; PF00132; hexapep; 3.
 SQ SEQUENCE 265 AA; 28346 MW; 0CE25212 CRC32;

Query Match 55.6%; Score 35; DB 2; Length 265;
 Best Local Similarity 57.1%; Pred. No. 32;
 Matches 8; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 1 YRLAIRXRILL 14
 |||||
 DB 40 YRLANRLWRAAWRY 53

RESULT 9
 ID Q57090 PRELIMINARY; PRT; 402 AA.
 AC Q57090;
 DT 01-NOV-1996 (TEMBLrel. 01, Created)
 DT 01-NOV-1996 (TEMBLrel. 01, Last sequence update)
 DT 01-NOV-1998 (TEMBLrel. 08, Last annotation update)
 DE TRANSPOSASE.
 OS Corynebacterium xerosis.
 OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
 OC Actinomycetales; Corynebacteriaceae; Corynebacterium.
 RN [1]

RP SEQUENCE FROM N.A.
 RC STRAIN-M82B;
 RX MEDLINE: 96117603.
 RA TAUCH A., KASSING F., KALINOWSKI J., PUHLER A.;
 RT "The Corynebacterium xerosis composite transposon Tn5432 consists of
 RT two identical insertion sequences, designated IS1249, flanking the
 RT erythromycin resistance gene ermCX.";
 RL Plasmid 34:119-131(1995).
 DR EMBL: U21300; AAC95477.1; -
 DR EMBL: U21300; AAC95474.1; -
 DR EMBL: U21300; AAC95474.1; -
 SQ SEQUENCE 402 AA; 45846 MW; 26339FD8 CRC32; -

Query Match 55.6%; Score 35; DB 2; Length 402;
 Best Local Similarity 60.0%; Pred. No. 49;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 1 YRLAIRIXRI 10
 |||::: ||
 Db 210 YRLAKLTRI 219

RESULT 10
 QXKIL2
 ID Q9XIL2 PRELIMINARY: PRT; 404 AA.
 AC Q9XIL2;
 DT 01-NOV-1999 (TRENBLrel. 12, Created)
 DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE PUTATIVE CYCLOPHILIN-TYPE PEPTIDYL-PROLYL CIS-TRANS ISOMERASE (EC
 DE 5.2.1.6)
 GN F19G14.21.
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC eudicotyledons; Spermatophyta; Magnoliophyta; eudicotyledons; core
 OC eudicotyledons; Rosidae; eurosids II; Brassicales; Brassicaceae;
 OC Arabidopsis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-CV. COLUMBIA;
 RA LIN X., KAUL S., SHEA T.P., FUJII C.Y., SHEN M., VANAKEN S.E.,
 RA BARNSTEAD M.E., MASON T.M., BOWMAN C.L., RONNING C.M., BENITO M.,
 RA CARRERA A.J., CREAMY T.H., BUELL C.R., TOWN C.D., NIERMAN W.C.,
 RA FRASER C.M., VENTER J.C.;
 RT "Arabidopsis thaliana chromosome II BAC F19G14 genomic sequence.";
 RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
 CC -1- CATALYTIC ACTIVITY: CIS-TRANS ISOMERIZATION OF PROLINE IMIDIC
 CC PEPTIDE BONDS IN OLIGOPEPTIDES.
 DR EMBL: AC006438; AAD41985.1; -
 DR PROSITE: PS00170; CSA_PP1ASE_1; 1.
 KW Isomerase; Rotamase.
 SQ SEQUENCE 404 AA; 45001 MW; A8A760CD CRC32;

Query Match 55.6%; Score 35; DB 10; Length 404;
 Best Local Similarity 50.0%; Pred. No. 49;
 Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRI 14
 ||::: |||
 Db 228 YKMLKRYKALRY 241

RESULT 11
 OS0159
 ID OS0159 PRELIMINARY: PRT; 469 AA.
 AC OS0159;
 DT 01-JUN-1998 (TRENBLrel. 06, Created)
 DT 01-JUN-1998 (TRENBLrel. 06, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE PROTON-TRANSLOCATING ATPASE, BETA SUBUNIT (EC 3.6.1.34).
 GN ATPD.
 OS Streptococcus bovis.

OC Bacteria; Firmicutes; Bacillus/Clostridium group; Streptococcaceae;
 OC Streptococcus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-JB-1;
 RA UMEMORI J., MIWA T., NAGAMINE T., OGATA K., TAKENAKA A., HINO T.;
 RL Submitted (NOV-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AB009314; BAA23755.1; -
 DR PROSITE: PS00152; ATPASE_ALPHA_BETA; 1.
 DR PFAM: PF00006; ATP-synt_ab; 1.
 DR PFAM: PF00306; ATP-synt_ab_C; 1.
 KW Hydrolase; Hydrogen ion transport.
 SQ SEQUENCE 469 AA; 51219 MW; A4170D3F CRC32;

Query Match 55.6%; Score 35; DB 2; Length 469;
 Best Local Similarity 42.9%; Pred. No. 57;
 Matches 6; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

QY 1 YRLAIRIXRI 14
 ||::: |||
 Db 362 YEVATEVQVQLQRY 375

RESULT 12
 O49025
 ID O49025 PRELIMINARY: PRT; 44 AA.
 AC O49025;
 DT 01-JUN-1998 (TRENBLrel. 06, Created)
 DT 01-JUN-1998 (TRENBLrel. 06, Last sequence update)
 DT 01-AUG-1998 (TRENBLrel. 07, Last annotation update)
 DE PLASMID GCH7220, COMPLETE SEQUENCE.
 OS Gracilaria chilensis.
 OC Plasmid Gch7220.
 OC Eukaryota; Rhodophyta; Florideophyceae; Gracilariaceae; Gracilariaceae;
 OC Gracilaria.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA GOFF L.J., MOON D.A.;
 RL Submitted (NOV-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL: AF034719; AAC04733.1; -
 KW Plasmid.
 SQ SEQUENCE 44 AA; 5487 MW; 6A285329 CRC32;

Query Match 54.0%; Score 34; DB 10; Length 44;
 Best Local Similarity 35.7%; Pred. No. 8.6;
 Matches 5; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY 1 YRLAIRIXRI 14
 ||::: |||
 Db 23 YRMCVKVKILYFY 36

RESULT 13
 Q9YC92
 ID Q9YC92 PRELIMINARY: PRT; 157 AA.
 AC Q9YC92;
 DT 01-NOV-1999 (TRENBLrel. 12, Created)
 DT 01-NOV-1999 (TRENBLrel. 12, Last sequence update)
 DT 01-NOV-1999 (TRENBLrel. 12, Last annotation update)
 DE 157AA LONG HYPOTHETICAL PROTEIN.
 GN APE1362.
 OS Aeropyrum pernix.
 OC Archaea; Crenarchaeota; Aeropyrum.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-K1;
 RX MEDLINE: 99310339.
 RA KANARABAYASI Y., HINO Y., HORIKAWA H., YAMAZAKI S., HAIKAWA Y.,
 RA JIN-NO K., TAKAHASHI M., SEKINE M., BABA S., ANKAI A., KOSUGI H.,
 RA HOSOIYAMA A., FUKUI S., NAGAI Y., NISHIJIMA K., NAKAZAWA H.,
 RA TAKAMIYA M., MASUDA S., FUNAHASHI T., TANAKA T., KUDOH Y.,

RA YAMAZAKI J., KUSHIDA N., OGUCHI A., AOKI K., KUBOTA K., NAKAMURA Y.,
RA NOMURA N., SAKO Y., KIKUCHI H.;
RT "Complete genome sequence of an aerobic hyper-thermophilic
RT crenarchaeon, Aeropyrum pernix K1.";
RL DNA Res. 6:83-101(1999)
DR EMBL: AP000061; BAA80356.1; -.
SQ SEQUENCE 157 AA; 17942 MW; C1F4AB62 CRC32;

Query Match 54.0%; Score 34; DB 1; Length 157;
Best Local Similarity 58.3%; Pred. No. 30;
Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 3 LAIRIXRILRY 14
||:| |||:|
DB 35 LAAKIARILTKY 46

RESULT 14

P94457
ID P94457 PRELIMINARY; PRT; 170 AA.
AC P94457;
DT 01-MAY-1997 (TReMBLrel. 03, Created)
DT 01-MAY-1997 (TReMBLrel. 03, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE ORF4 GENE (FRAGMENT).
OS Bacillus stearothermophilus.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Bacillus.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-NCIB 8224;
RX MEDLINE: 94113715.
RA SAKANYAN V., DESMAREZ L., LEGRAIN C., CHARLIER D., METT T.,
RA KOCHIKYAN A., SAVCHENKO A., BOYEN A., FALMAGNE P., PIRARD A.,
RA GLANDORFF N.;
RT "Gene cloning, sequence analysis, purification, and characterization
RT of a thermostable aminoacylase from Bacillus stearothermophilus.";
RL Appl. Environ. Microbiol. 59:3878-3888(1993).
DR EMBL: Y08751; CAA69998.1; -.
FT NON_TER 1
SQ SEQUENCE 170 AA; 18433 MW; 60A9C69B CRC32;

Query Match 54.0%; Score 34; DB 2; Length 170;
Best Local Similarity 50.0%; Pred. No. 33;
Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILRY 14
|||:| |||
DB 32 YRLEMTTEKLLRY 45

RESULT 15

Q9YFDO
ID Q9YFDO PRELIMINARY; PRT; 276 AA.
AC Q9YFDO;
DT 01-NOV-1999 (TReMBLrel. 12, Created)
DT 01-NOV-1999 (TReMBLrel. 12, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE 276AA LONG HYPOTHETICAL CITRATE LYASE BETA CHAIN.
GN APE0311.
OS Aeropyrum pernix.
OC Archaea; Crenarchaeota; Aeropyrum.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-K1;
RX MEDLINE: 99310339.
RA KAWARABAYASI Y., HINO Y., HORIKAWA H., YAMAZAKI S., HAIKAWA Y.,
RA JIN-NO K., TAKAHASHI M., SEKINE M., BABA S., ANKAI A., KOSUGI H.,
RA HOSOTAYAMA A., FUKUI S., NAGAI Y., NISHIJIMA K., NAKAZAWA H.,
RA TAKAMIYA M., MASUDA S., FUNAHASHI T., TANAKA T., KUDOH Y.,
RA YAMAZAKI J., KUSHIDA N., OGUCHI A., AOKI K., KUBOTA K., NAKAMURA Y.,

RA NOMURA N., SAKO Y., KIKUCHI H.;
RT "Complete genome sequence of an aerobic hyper-thermophilic
RT crenarchaeon, Aeropyrum pernix K1.";
RL DNA Res. 6:83-101(1999)
DR EMBL: AP000059; BAA79266.1; -.
KW Lyase.
SQ SEQUENCE 276 AA; 30725 MW; 4B749B21 CRC32;

Query Match 54.0%; Score 34; DB 1; Length 276;
Best Local Similarity 50.0%; Pred. No. 53;
Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 YRLAIRIXRILRY 14
|||:| |||
DB 259 YRLALNLLRRASY 272

Search completed: February 8, 2000, 13:17:49
Job time: 32498 sec

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Date: Feb 8, 2000 4:44 PM

About: Results were produced by the GenCore software, version 4.5,
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Command line parameters:

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-MINMATCH=0.100 -LOPCU=0.000 -LOPEXT=0.000 -CGAPOP=4.500
-CGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500 -FGAPOP=6.000
-FGAPEXT=7.000 -GAPOP=10.000 -XGAPEXT=0.500 -DELOP=6.000
-DELEXT=7.000 -START=1 -MATRIX=biosum62 -TRANS=human40.cdi
-LIST=45 -LOCALIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
-OUTFMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=1000000 -USER=US08653294
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Search information block:

Query: US-08-653-294-20
Query length: 14
Database: GenEmbl:*
Database sequences: 821193
Database length: 1518192014
Search time (sec): 11370.480000

score_list:

Sequence	Strd Orig	ZScore	EScore	Len	Documentation
gb_ro:AC002121	-	43.00	101.77	84056	AC002121 Genomic sequence from Mouse 11
gb_ba1:AP089957	+	42.00	120.19	113.36	U89957 Actinobacillus pleuropneumoniae
gb_bt2:AP000657	+	42.00	102.43	50660	AP000657 Homo sapiens genomic clone
gb_bt6:AC013287	+	42.00	91.66	172450	AC013287 Homo sapiens clone
gb_p11:SCYPL167C	-	41.00	119.30	127.11	Z73523 S.cerevisiae chromosome
gb_p11:HUMOPIDRE	+	41.00	119.28	4839	M4605 Human putative opitoid
gb_p11:YSGREV3	+	41.00	118.90	123.43	M29683 S.cerevisiae DNA polym
gb_p11:SCACHXVI	-	41.00	97.79	55786	X96770 S.cerevisiae chromosome
gb_bt5:AC013211	+	40.00	117.79	154.41	AC013211 Drosophila melanogaster
gb_in1:CE802A10	+	40.00	101.45	23889	Z81053 Caenorhabditis elegans
gb_in1:CEW01C1	+	40.00	97.76	2.0e+03	Z46381 Caenorhabditis elegans
gb_in1:CEZ21C9	+	40.00	97.48	36355	Z73098 Caenorhabditis elegans
gb_p12:AB028622	-	40.00	90.57	82348	AB028622 Arabidopsis thaliana
gb_p12:ATAC007070	-	40.00	90.19	86017	AC007070 Arabidopsis thaliana
gb_bt2:AC009548	+	40.00	88.89	99657	AC009548 Homo sapiens chromosome
gb_bt1:CEV2A483	+	40.00	86.11	136764	AL008873 Caenorhabditis elegans
gb_p12:AC002456	+	40.00	84.93	156399	AC002456 Human BAC clone RQ
gb_bt2:AC013355	+	40.00	83.17	191137	AL133355 Homo sapiens chromosome
gb_bt6:AC012517	+	40.00	81.39	233904	AC012517 Homo sapiens clone
gb_p12:ABU54809	-	39.00	128.15	745	U54809 Armillaria borealis rDNA
gb_p13:AF056334	+	39.00	112.81	4265	AF056334 Homo sapiens cancer
gb_ba1:MVOMCR	-	39.00	110.71	5412	M16893 Methanococcus vannielii
gb_p13:AF064589	+	39.00	109.14	6471	X0764589 Homo sapiens melanoma
gb_ba1:BSVWCR1	-	39.00	109.14	6472	X07793 Methanococcus voltae
gb_ba1:BSVWCR1	-	39.00	107.57	7735	Z35133 B.subtilis 168 pks gene
gb_bt7:AC018125	+	39.00	106.13	9116	AC018125 Drosophila melanogaster
gb_bt7:AC017410	+	39.00	95.72	29776	AC017410 Drosophila melanogaster
gb_bt2:AC006614	-	39.00	94.61	33803	AC006614 Caenorhabditis elegans
gb_bt5:AC015441	+	39.00	94.46	34357	AC015441 Drosophila melanogaster
gb_ba1:MTY373	+	39.00	94.17	35516	Z73419 Mycobacterium tuberculosis
gb_ba1:MLU15186	+	39.00	93.99	36241	U15186 Mycobacterium leprae
gb_in2:AC005447	+	39.00	87.27	77907	AC005447 Drosophila melanogaster
gb_p12:ATAC007187	+	39.00	85.21	98432	AC007187 Arabidopsis thaliana
gb_p12:ATF4110	+	39.00	84.08	111876	AL035525 Arabidopsis thaliana
gb_p12:CN5010RM	+	39.00	83.60	118261	AL117687 Human chromosome 14
gb_bt2:CN508259	+	39.00	83.49	119692	AC008259 Drosophila melanogaster
gb_p13:HS326112	-	39.00	83.44	120423	AL023279 Homo sapiens DNA se
gb_p13:HS326224	-	39.00	83.14	124497	AL022152 Homo sapiens DNA se
gb_bt3:HS232624	+	39.00	82.92	127727	AC009393 Drosophila melanogaster
gb_bt3:AC009393	+	39.00	82.50	133889	AC031643 Human DNA sequence
gb_p13:HS76920	+	39.00	80.92	160359	AC007262 Homo sapiens chromosome
gb_p13:AC007262	+	39.00	80.30	172043	AC011614 Drosophila melanogaster

gb_pr2:CN50000E - 39.00 79.54 2.1e+04 187564 ! AL049835 Human chromosome
gb_ba1:BSUB0010 + 39.00 77.60 2.6e+04 233780 ! Z99113 Bacillus subtilis
gb_p11:AB017564 + 38.00 121.92 90.89 983 ! AB017564 Arabidopsis thaliana

seq_name: gb_ro:AC002121

seq_documentation_block:

LOCUS AC002121 84056 bp DNA 10-JUL-1997
DEFINITION Genomic sequence from Mouse 11, complete sequence.

ACCESSION AC002121

VERSION AC002121.1 GI:2133880

KEYWORDS HTG.

SOURCE house mouse.

ORGANISM Mus musculus

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

REFERENCE 1 (bases 1 to 84056)

Hawkins,T.L., Reeve,M.P., Christoffersen,A., Birren,B.W.,
Fasman,K.H. and Lander,E.S.

Unpublished
Genomic sequence from Mouse 11

REFERENCE 2 (bases 1 to 84056)

Hawkins,T.L., Reeve,M.P., Christoffersen,A., Birren,B.W.,
Fasman,K.H., Lander,E.S., McKernan,K., Munro,C., Richardson,P.,
Barna,N., Chang,A., Cooke,P., Daly,M.J., Forrest,C., Frapp,W.J.,
Gage,D., Geraghty,K., Hagos,B., Jacotot,L., Lane,M., Mackenzie,J.,
Marquis,N., McDermott,J., Moloney,N., Morrow,J., Nachman,A.,
Naylor,J., O'Connor,T., Peterson,K., Rollins,G., Spencer,J.,
Stiwell,J., Stone,C., Strickland,C., Sydney,K., Wilmer,F. and
Zody,M.

Direct Submission

Submitted (14-MAY-1997) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

REFERENCE 3 (bases 1 to 84056)

Hawkins,T.L., Reeve,M.P., Christoffersen,A., Birren,B.W.,
Fasman,K.H., Lander,E.S., McKernan,K., Munro,C., Richardson,P.,
Barna,N., Chang,A., Cooke,P., Daly,M.J., Forrest,C., Frapp,W.J.,
Gage,D., Geraghty,K., Hagos,B., Jacotot,L., Lane,M., Mackenzie,J.,
Marquis,N., McDermott,J., Moloney,N., Morrow,J., Nachman,A.,
Naylor,J., O'Connor,T., Peterson,K., Rollins,G., Spencer,J.,
Stiwell,J., Stone,C., Strickland,C., Sydney,K., Wilmer,F. and
Zody,M.

Direct Submission

Submitted (29-MAY-1997) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

REFERENCE 4 (bases 1 to 84056)

Hawkins,T.L., Reeve,M.P., Christoffersen,A., Birren,B.W.,
Fasman,K.H., Lander,E.S., McKernan,K., Munro,C., Richardson,P.,
Barna,N., Chang,A., Cooke,P., Daly,M.J., Forrest,C., Frapp,W.J.,
Gage,D., Geraghty,K., Hagos,B., Jacotot,L., Lane,M., Mackenzie,J.,
Marquis,N., McDermott,J., Moloney,N., Morrow,J., Nachman,A.,
Naylor,J., O'Connor,T., Peterson,K., Rollins,G., Spencer,J.,
Stiwell,J., Stone,C., Strickland,C., Sydney,K., Wilmer,F. and
Zody,M.

Direct Submission

Submitted (10-JUL-1997) Whitehead Institute/MIT Center for Genome

Research, 320 Charles Street, Cambridge, MA 02141, USA

COMMENT

On May 29, 1997 this sequence version replaced gi:2098549.

The Staden databases, finishing information, and all

chromatographic files used in the assembly of this clone are

available from our anonymous ftp site.

All repeats were identified using RepeatMasker: Smit, A.F.A. &
Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html.

Location/Qualifiers

1. 84056

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/clone="5157"

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complement(1051..1109)
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6842..6988
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12840..12994
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13021..13139
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  Ratio: 3.583      Gaps: 0
  Percent Similarity: 85.714      Percent Identity: 64.286

alignment_block:
US-08-653-294-20 x AC002121/rev

Align seg 1/1 to reverse of: AC002121 from: 1 to: 84056

1 TyrArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14

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[illegible]

KRHVSGHQWQSSGEFEAFYKK

CDS 94. .1416

94. .1416

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PRGPAPHPFQPPMAVALWSLAYGAVAVAVGLNVLVIVILAHKRMRTVITNSFLVNL
AFADAAAMALNVLNVIYALHGEWYFGANYCRFNFFPTAVFASITYTAIADRYM
AIDIDLPRLSATRTIVIGSIWILAFPOCLYKIKVMPGRTLCVQWPEGSRQ
HFTYHVIIVLVVCEPLLINGITVITVGLWGEIPGDCDYQOLAKAKRVKWM
IIVVFAICWLPYHLYFILTALYQOLNWKYIQOYVLASFWLAMSSTWNPILYCL
NKRPRAGFRKAWCFHVSYSDELELKATIRLHPKRSQSLYTVTRHMSVVFDSND
GDSRASHQKRGRTTRDVGNSRRRSKSTSTTASFVSSHMSVEEGS"
BASE COUNT      1449 a   1099 c   965 g   1326 t
ORIGIN

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alignment_scores:
  Quality: 41.00      Length: 13
  Ratio: 3.417        Gaps: 0
  Percent Similarity: 92.308  Percent Identity: 76.923

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alignment_block:
US-08-653-294-20 x HUMOPIOORE ..

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Align seg 1/1 to: HUMOPIOORE from: 1 to: 4839

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```

1  ArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
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1530 AGATTGCATATAATATACAAATACTACTAGATAT 1568

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seq_name: gb_pl1:YSCREV3

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seq_documentation_block:
LOCUS      YSCREV3      5056 bp      DNA      PLN      27-APR-1993
DEFINITION YSCREV3 Saccharomyces cerevisiae (rev3) gene, complete cds.
ACCESSION  M29683
VERSION    M29683.1 GI:172386
KEYWORDS   DNA polymerase.
SOURCE     Saccharomyces cerevisiae
ORGANISM   Eukaryota; Fungi; Ascomycota; Saccharomycetales;
            Saccharomycetaceae; Saccharomycetes.
REFERENCE  Morrison,A., Christensen,R.B., Alley,J., Beck,A.K., Bernstine,E.G.,
            Lemontt,J.F. and Lawrence,C.W.
            REV3, a Saccharomyces cerevisiae gene whose function is required
            for induced mutagenesis, is predicted to encode a nonessential DNA
            polymerase
JOURNAL    J. Bacteriol. 171, 5659-5667 (1989)
MEDLINE    90008808
FEATURES   Location/Qualifiers
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              /organism="Saccharomyces cerevisiae"
              /db_xref="taxon:4932"
              305..4819
              /note="DNA polymerase (pot.): putative"
              /codon_start=1
              /db_xref="GI:172387"
              /protein_id="AAA34968.1"
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            ISLNPSCSLSELIRDGKIFGKFEIYESHPIYLLQWTFADNFGCSQVNVDRCYF
            RSPVLNSILDILTDLQLLDRFCDFKCNVLSRRDFPRVNGLIETDILPQFIK
            NREKLOHRDHFLEKLDISDIPVKVYSARDMINELTQRELSLKEVKEPPT
            KRVSGHOMSGSEFEAFYKKAHKISTDGCQIPNFENIDNKNKESAINPYEALPQ
            LWPRLPOIELNNMQDKNDQVNASFTEIECGVDNENGVKNRSKRSYSWLPQ
            SIAPKSDTLLDQTKYHNTINFSDCANTONMASKRLRSVSANKTSLSRKRYK
            VMAAGLYGRARVYGPFGYQDILNKLDEGFPKIDYKDPFSNPVDLENKPYAY
            GKRFISSTHTRIPVQFGGETVSVYKPTFDMSSWKYALKPPTYDAVQKYNKVP
            SMGNKTESQISMHTSPKFLYKFAFSDVSKOKRKSVDHSLTTLTLEHANTRSK
            IPPDADEVSMITWCEEFPLDLDIAYEGIMVHKASDSTFFPKIQCINCIEIPVM
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```

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VLDOSLYPSIMIGNYCYSTMIGRVREINLTENNLSGSKFSLPRLNALLKNDVTIA
PNGVYAKTSVSKSLSKMLTDILOVRVNIKKTMNEIGDDNTTLRLNKKQLAKLL
ANYTGITSAFSGMRPCSDLADSVOTGRELEKAIDIEKDETWNKAVVIGDTDSL
FYLPGKTAIEAFSGHAMAERTQNNPKPIFLKFEKVYHPSILISKRYVGSFYESP
SOTLIFDAKGIETVRDGIQAQKIIEKIRLLFTQDKLSIKKYLQNEFFQIQIGK
VSAQDFCAKEVKGAYKSEKAPAGAVVVKRRINEDHRAEQYKERIPYLVVKGQK
QILLRECVSEPEELEGLELDSEYVINKILIPDLDRLENLGINVGNWAOETVKSRR
ASTTTKVENIIRVGTSAATCCNGBELFKISQILQDCGLEKRSITTLFLKLLKRRQ
KEYTTLKTCRTSRITSAGIENHASKNSYDCPVFSRVAERYLRDNQSVQR
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BASE COUNT      1770 a   856 c   972 g   1458 t
ORIGIN

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alignment_scores:
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  Ratio: 3.727        Gaps: 0
  Percent Similarity: 78.571  Percent Identity: 57.143

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alignment_block:
US-08-653-294-20 x YSCREV3 ..

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Align seg 1/1 to: YSCREV3 from: 1 to: 5056

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1  TyrArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
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1647 TACCGGAAGCATTCGTCCTCCGAAGATCTACTATAT 1688

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seq_name: gb_pl1:SCLACHXVI

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seq_documentation_block:
LOCUS      SCLACHXVI      55786 bp      DNA      PLN      03-DEC-1996
DEFINITION S.cerevisiae chromosome XVI, left arm DNA.
ACCESSION  X96770
VERSION    X96770.1 GI:1409537
KEYWORDS   BEM4 gene; CDC60 gene; KES1 gene; KIP2 gene; OYE3 gene; PAL1 gene;
            PEP4 gene; PXA1 gene; REV3 gene; ribosomal protein L37a; RPL37A
            gene; SNR178 gene; spk1 gene; SVS1 gene; u3 small nuclear rna.
            baker's yeast.
SOURCE     Saccharomyces cerevisiae
ORGANISM   Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
            Saccharomycetaceae; Saccharomycetes.
REFERENCE  1 (bases 1 to 55786)
            Purnelle,B., Coster,F. and Goffeau,A.
            The sequence of 55 kb on the left arm of yeast chromosome XVI
            identifies a small nuclear RNA, a new putative protein kinase and
            two new putative regulators
JOURNAL    Yeast 12 (14), 1483-1492 (1996)
MEDLINE    97103777
FEATURES   Location/Qualifiers
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              /map="left arm"
            /direct_submission
            Submitted (22-MAR-1996) B. Purnelle, Unite de Biochimie
            Physiologique, Universite Catholique de Louvain, Place Croix du Sud
            2/20, 1348 Louvain-la-Neuve, BELGIUM
            Overlapping sequences: L29279, M29683, D50278, L27816, X62878,
            211963, M1358, X55623, L38491, U17065, U03913, X57969, X05498.
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JOURNAL    Direct Submission
MEDLINE    97103777
REFERENCE  2 (bases 1 to 55786)
            Purnelle,B.
            The sequence of 55 kb on the left arm of yeast chromosome XVI
            identifies a small nuclear RNA, a new putative protein kinase and
            two new putative regulators
JOURNAL    Yeast 12 (14), 1483-1492 (1996)
MEDLINE    97103777
FEATURES   Location/Qualifiers
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            /clone="cos8484"
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1  T+ArgLeuAlaIleArgIle**ArgIleLeuLeuArgTyr 14
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3148 TATAGCATAGCTGAAGATTCGAAAAATGCTACTTCGATAT 3189

seq name: qb in1:CEE02A10

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REFERENCE
1 (bases 1 to 23889)
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
Zaruafoa; rotaria; nematoda; rhabditidae; peloderinae; caenorhabditis.

Fulton, L., Gardner, A., Green, P., Hawkins, T., Hillier, L., Jier, M., Johnston, L., Jones, M., Kershaw, J., Kirsten, J., Laister, N.,

O'Callaghan, M., Parsons, J., Percy, C., Rifken, L., Roopra, A.,

Staden, R., Sulston, J., Thierry-Mieg, J., Thomas, K., Vaudin, M., Vaughan, K., Waterston, R., Watson, A., Weinstock, L., Wilkinson-Sproat, J., and Woldman, P.

JOURNAL Nature 368 (6456), 32-38 (1994)
MEDLINE 94150718
REFERENCE 2 (bases 1 to 23889)
AUTHORS Thomas, K.

Submitted (21-OCT-1996) Louis, MO 63110, USA. E-mail:
jestsanger.ac.uk or rwnematode.wustl.edu
On Dec 12, 1996 this sequence version replaced gi:1627715.

predictions from genefinder (P. Green, U. Washington), and other available information.
For a graphical representation of this sequence and its analysis see:-

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bin/display?db=wormacsclass=Sequence object=E02A10
http://www.ncbi.nlm.nih.gov/seq/
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dye primer and dye terminator' reaction, from distinct subclones. Exceptions are indicated by an explicit note.

IMPORTANT: This sequence is NOT necessarily the entire insert of

overlapping sections once, or longer because we arrange for a small overlap between neighbouring submissions.

once, or longer because we arrange for a small overlap between neighbouring submissions.

The true left end of clone C14C10 is at 23796 in this sequence. The true right end of clone F45D3 is at 107 in this sequence. The start

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FEATURES
source
      Location/Qualifiers
      1. .23889
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         /db_xref="taxon:6239"
sequence Z74028.
The end of this sequence (23793..23889) overlaps with the start of
the next sequence (23889..23900)

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gene
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complement(11007..12594)
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complement(join(11007. .11504,11551. .11813,11984. .12016,
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seq_documentation_block:
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DEFINITION  Caenorhabditis elegans cosmid M01F1, complete sequence.
ACCESSION   Z46381
VERSION     246381.1  GI:561920
KEYWORDS    HTG; Glucose transport protein; L13P ribosomal protein; Lipoic acid
            synthase; ysy6.

SOURCE      Caenorhabditis elegans.
ORGANISM    Caenorhabditis elegans
            Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
            Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
            1 (bases 1 to 36355)

REFERENCE   1 (bases 1 to 36355)
AUTHORS    Wilson, R., Ainscough, R., Anderson, K., Baynes, C., Berks, M.,
            Bonfield, J., Burton, J., Connell, M., Copsey, T., Cooper, J.,
            Coulson, A., Craxton, M., Dear, S., Du, Z., Durbin, R., Favello, A.,
            Fulton, L., Gardner, A., Green, P., Hawkins, T., Hillier, L., Jier, M.,
            Johnston, L., Jones, M., Kershaw, J., Kirsten, J., Laister, N.,
            Latreille, P., Lightning, J., Lloyd, C., McMurray, A., Mortimore, B.,
            O'Callaghan, M., Parsons, R., Percy, C., Rifken, L., Roopra, A.,
            Saunders, D., Showkneen, R., Smaldon, N., Smith, A., Sonhammer, E.,
            Vaughan, R., Sulston, J., Thierry-Mieg, J., Thomas, K., Vaudin, M.,
            Vaughan, K., Waterston, R., Watson, A., Weinstein, L.,
            Wilkinson-Spratt, J. and Wohldman, P.
            2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
            elegans

JOURNAL     Nature 368 (6466), 32-38 (1994)
MEDLINE     94150718
REFERENCE   2 (bases 1 to 36355)
AUTHORS     Sims, M.
            Direct Submission
            Submitted (29-OCT-1994)  Louis, MO 63110, USA.  E-mail:
            jess@sanger.ac.uk or rw@nematode.wustl.edu
            Coding sequences below are predicted from computer analysis, using
            predictions from Gensfinder (P. Green, U. Washington), and other
            available information.
            For a graphical representation of this sequence and its analysis
            see:
            http://webace.sanger.ac.uk/cgi-
            bin/display?db=wormacc&class=Sequence&object=M01F1
            Current sequence finishing criteria for the C. elegans genome
            sequencing consortium are that all bases are either sequenced
            unambiguously on both strands, or on a single strand with both a
            dye primer and dye terminator reaction, from distinct subclones.
            Exceptions are indicated by an explicit note.
            IMPORTANT: This sequence is NOT necessarily the entire insert of
            the specified clone. It may be shorter because we only sequence
            overlapping sections once, or longer because we arrange for a small
            overlap between neighbouring submissions.
            IMPORTANT: This sequence is not the entire insert of clone M01F1.
            It may be shorter because we only sequence overlapping sections
            once, or longer because we arrange for a small overlap between
            neighbouring submissions.
            The true left end of clone M01F1 is at 1 in this sequence. The
            start of this sequence (1..104) overlaps with the end of sequence
            Z46396.
            The end of this sequence (36252..36355) overlaps with the start of
            sequence Z77131.

FEATURES             Location/Qualifiers
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                     /clone="M01F1"
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                     5651..5706,5757..5807,6111..6251,6299..6382))
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                     /note="Similar to alpha-1,3-mannosyl-glycoprotein
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NLQKFKPIYISRHKALNHLIFSNSNYSSVIIIEDLDIADDFISFISNRIILEK
DPSLMCVTAWNDGKPNIDLSKNATLYRSDFFAGLGMWMTKRTEWEELEWPNFWD
DMWRFPVQRQCRPEISRTGMKYGKESQKQFSDHLEKIKVNDLVPDFSQIN
LDYLQKNEFESRLSDIRNAPVDIDDIYTPDMKPDYEGMKAIYITGRTDFVAKDR
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15281..15487,15533..15694))
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/notes="cDNA EST yk415a9.3 comes from this gene; cDNA EST
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from this gene"
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FYAREQAPNERGVCWRAALADEYHDIADIVNAGROTIVRIPKAGVCPAPLQORP
LAPPNPQVPSSTANNIAITPTPTVTQRLRPVTPQPVKSFAQAKPPIDGDFYDFS
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gene; cDNA EST EMBL:D72496 comes from this gene; cDNA EST
EMBL:D68294 comes from this gene; cDNA EST EMBL:D73026
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gene; cDNA EST EMBL:D75395 comes from this gene; cDNA EST
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comes from this gene; cDNA EST EMBL:D70026 comes from this
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EMBL:C09924 comes from this gene; cDNA EST EMBL:C08172
comes from this gene; cDNA EST EMBL:C07790 comes from this
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gene; cDNA EST EMBL:C08961 comes from this gene; cDNA EST
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comes from this gene; cDNA EST yk503e5.3 comes from this
gene; cDNA EST yk500g9.3 comes from this gene; cDNA EST
yk486g2.3 comes from this gene; cDNA EST yk486g2.5 comes
from this gene; cDNA EST yk475e7.3 comes from this gene;
cDNA EST yk475e7.5 comes from this gene; cDNA EST
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from this gene; cDNA EST yk400a3.3 comes from this gene;
cDNA EST yk400a3.5 comes from this gene; cDNA EST
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from this gene; cDNA EST yk372b4.3 comes from this gene;
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cDNA EST yk298a4.5 comes from this gene; cDNA EST
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from this gene; cDNA EST yk327c5.3 comes from this gene;
cDNA EST yk327c5.5 comes from this gene; cDNA EST
yk300b7.3 comes from this gene; cDNA EST yk300b7.5 comes
from this gene; cDNA EST yk313a3.3 comes from this gene;
cDNA EST yk313a3.5 comes from this gene; cDNA EST
yk252e3.3 comes from this gene; cDNA EST yk252e3.5 comes
from this gene; cDNA EST yk235h1.3 comes from this gene;
cDNA EST yk235h1.5 comes from this gene; cDNA EST
yk235b11.3 comes from this gene; cDNA EST yk235b11.5 comes
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complement(16937..18194)
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CDS
complement(join(16937..17265,17322..17557,17602..17903,
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comes from this gene; cDNA EST yk283b6.5 comes from this
gene; cDNA EST yk472f5.3 comes from this gene; cDNA EST
yk472f5.5 comes from this gene; cDNA EST yk476e7.3 comes
from this gene; cDNA EST yk476e7.5 comes from this gene;
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Ratio:	3.636	Gaps:	0
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US-08-653-294-20 x CEM01F1/rev ..

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1 TyrArgLeuAlaIleArgIle***ArgIleLeuLeuArg 13

4482 TTGAGCTGAAATTCGAATTCGCGCATTTGTACGT 4444

seq_name: gb_inl:CET21C9

seq_documentation_block:

[illegible]

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Percent Similarity: 71.429 Percent Identity: 57.143

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US-08-653-294-20 x CET21C9 ..
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seq_name: gb_pl1:AB028622
seq_documentation_block:
LOCUS AB028622 82348 bp DNA PLN 20-NOV-1999

DEFINITION Arabidopsis thaliana genomic DNA, chromosome 3, P1 clone:MZN24, complete sequence.
ACCESSION AB028622
VERSION AB028622.1 GI:5041975
KEYWORDS HTG
SOURCE Arabidopsis thaliana (strain:Columbia) DNA, clone_lib:Mitsui P1 clone:MZN24.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsi.
REFERENCE 1 (sites)
AUTHORS Nakamura,Y.
TITLE Structural Analysis of Arabidopsis thaliana Chromosome 3. II
JOURNAL Unpublished (1999)
REFERENCE 2 (bases 1 to 82348)
AUTHORS Nakamura,Y.
TITLE Direct Submission
JOURNAL Submitted (09-JUN-1999) to the DDBJ/EMBL/GenBank databases. Yasukazu Nakamura, Kazusa DNA Research Institute, Laboratory of Gene Structure 2; 1532-3, Yana, Kisarazu, Chiba 292, Japan (E-mail:ynakamu@kazusa.or.jp, Tel:++81-438-52-3935, Fax:++81-438-52-3934)
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BASE COUNT 27062 a 13968 c 13880 g 27438 t
ORIGIN

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Quality: 40.00 Length: 14
Ratio: 4.000 Gaps: 0
Percent Similarity: 71.429 Percent Identity: 50.000

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US-08-653-294-20 x AB028622 ..
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seq_documentation_block:
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DEFINITION Arabidopsis thaliana chromosome II BAC T22F11 genomic sequence, complete sequence.
ACCESSION AC007070
VERSION AC007070.3 GI:4567237
KEYWORDS HTG
SOURCE thale cress.
ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae; Arabidopsis.
REFERENCE 1 (bases 1 to 86017)
AUTHORS Lin,X., Kaul,S., Shea,T.P., Fujii,C.Y., Shen,M., VanAken,S.E., Barnstead,M.E., Mason,T.M., Bowman,C.L., Ronning,C.M., Benito,M., Carrera,A.J., Creasy,T.H., Buell,C.R., Town,C.D., Nierman,W.C., Fraser,C.M. and Venter,J.C.
TITLE Arabidopsis thaliana chromosome II BAC T22F11 genomic sequence
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 86017)


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*      gap of unknown length
* 19793 20604: contig of 812 bp in length
*      gap of unknown length
* 20605 21420: contig of 816 bp in length
*      gap of unknown length
* 21421 22242: contig of 822 bp in length
*      gap of unknown length
* 22243 23048: contig of 806 bp in length
*      gap of unknown length
* 23049 23890: contig of 842 bp in length
*      gap of unknown length
* 23891 24707: contig of 817 bp in length
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* 24708 25523: contig of 816 bp in length
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* 25524 26336: contig of 813 bp in length
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* 26337 27148: contig of 812 bp in length
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* 27957 28761: contig of 805 bp in length
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* 31178 31982: contig of 805 bp in length
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* 31983 32791: contig of 809 bp in length
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* 32792 33626: contig of 835 bp in length
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* 33627 34496: contig of 870 bp in length
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* 34497 35333: contig of 837 bp in length
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* 36801 37601: contig of 801 bp in length
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* 37602 38406: contig of 805 bp in length
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* 38407 39214: contig of 808 bp in length
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* 39215 40038: contig of 824 bp in length
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* 40039 40945: contig of 907 bp in length
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* 40946 41921: contig of 976 bp in length
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* 41922 42857: contig of 936 bp in length
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* 42858 43829: contig of 972 bp in length
*      gap of unknown length
* 43830 44705: contig of 876 bp in length
*      gap of unknown length
* 44706 45648: contig of 943 bp in length
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* 45649 46524: contig of 876 bp in length
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* 46525 47465: contig of 941 bp in length
*      gap of unknown length
* 47466 48319: contig of 854 bp in length
*      gap of unknown length
* 48320 49249: contig of 930 bp in length
*      gap of unknown length
* 49250 50175: contig of 926 bp in length
*      gap of unknown length
* 50176 51105: contig of 930 bp in length
*      gap of unknown length

* 51106 52091: gap of unknown length
*      contig of 986 bp in length
* 52092 53040: gap of unknown length
*      contig of 949 bp in length
* 53041 53966: gap of unknown length
*      contig of 926 bp in length
* 53967 54835: gap of unknown length
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* 54836 55789: gap of unknown length
*      contig of 954 bp in length
* 55790 56674: gap of unknown length
*      contig of 885 bp in length
* 56675 57631: gap of unknown length
*      contig of 957 bp in length
* 57632 58514: gap of unknown length
*      contig of 883 bp in length
* 58515 59397: gap of unknown length
*      contig of 883 bp in length
* 59398 60274: gap of unknown length
*      contig of 877 bp in length
* 60275 61163: gap of unknown length
*      contig of 889 bp in length
* 61164 62071: gap of unknown length
*      contig of 908 bp in length
* 62072 62952: gap of unknown length
*      contig of 881 bp in length
* 62953 63827: gap of unknown length
*      contig of 875 bp in length
* 63828 64798: gap of unknown length
*      contig of 971 bp in length
* 64799 65764: gap of unknown length
*      contig of 966 bp in length

alignment_scores:
  Quality: 40.00      Length: 13
  Ratio: 4.000       Gaps: 0
  Percent Similarity: 76.923  Percent Identity: 61.538

alignment_block:
US-08-653-294-20 x AC009548 ..
Align seg 1/1 to: AC009548 from: 1 to: 99657
      1 TyrArgLeuAlaIleArgIle***ArgIleLeuArg 13
      |||||
      96875 TACCGCCTTCCGCAAGGCGAGACCCGAGTAGTGTGCGC 96913
```

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OM of: US-08-653-294-20 to: N_Geneseq_36.* out_format : pfs

Date: Feb 8, 2000 1:28 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

```
-MODEL=frame+p2n.model -DEV=xlpl  
-O=/cgn1.1/USPTO_spool/US08653294/runat_04022000_160701_15807/app_query.fasta.1  
-DB=N_Geneseq_36 -OFMT=fastap -SUFFIX=tns -GAPOP=12.000  
-GAPEXT=4.000 -MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000  
-GAPOP=4.500 -GAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500  
-XGAPOP=6.000 -XGAPEXT=7.000 -XGAPOP=10.000 -XGAPEXT=0.500  
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=blotsum62  
-TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR_SCORE=pct  
-ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -MINLEN=0  
-MAXLEN=1000000 -USER=US08653294 -NCPU=6 -ICPU=3 -NO_XLPXY -WAIT  
-THREADS=1
```

Search information block:

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Query: US-08-653-294-20  
Query length: 14  
Database: N_Geneseq_36.*  
Database sequences: 311585  
Database length: 125096042  
Search time (sec): 590.520000
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score_list:

Sequence	Strd	Orig	Zscore	Escore	Len	Documentation
N_Geneseq_36:T84234	+	40.00	115.48	78.45	3110	DNA encoding an autolysin and
N_Geneseq_36:V53557	+	40.00	115.48	78.45	3110	DNA encoding 2 Staphylococcus
N_Geneseq_36:V74331	+	40.00	104.87	305.85	10470	Staphylococcus aureus contig
N_Geneseq_36:V69717	+	39.00	109.69	164.86	4031	Tumour rejection antigen precu
N_Geneseq_36:V69720	+	39.00	109.28	173.78	4225	Tumour rejection antigen precu
N_Geneseq_36:T48660	+	37.00	135.04	6.39	99	DNA encoding MAB anti-HSag bind
N_Geneseq_36:X24237	+	37.00	121.63	35.64	459	WO9916900 Seq ID 7. Identifying
N_Geneseq_36:X24233	+	37.00	121.63	35.64	459	WO9916900 Seq ID 7. Identifying
N_Geneseq_36:T33641	+	37.00	113.13	106.12	1215	Aspergillus arabinofuranosidase
N_Geneseq_36:Q98535	+	37.00	109.81	162.40	1776	Coding sequence for alkaline l
N_Geneseq_36:T51339	+	37.00	108.31	196.90	2109	Equine IFN-omega-1 from PAH61.
N_Geneseq_36:T33646	+	37.00	106.63	244.14	2555	Aspergillus arabinofuranosidase
N_Geneseq_36:N60309	+	37.00	108.31	196.90	2109	Equine IFN-omega-1 from PAH61.
N_Geneseq_36:V29571	+	37.00	101.09	496.71	4815	L. lactis soluble part (Fl) ex
N_Geneseq_36:V29571	+	37.00	98.16	723.54	6735	Streptococcus pneumoniae genom
N_Geneseq_36:X20568	+	37.00	96.79	862.03	7874	Polynucleotide sequence from t
N_Geneseq_36:X13331	+	37.00	91.68	1.7e+03	14141	Enterococcus faecalis genome
N_Geneseq_36:X13238	+	37.00	89.68	2.1e+03	17764	Enterococcus faecalis genome
N_Geneseq_36:T61973	+	36.00	103.90	346.44	2333	Pseudomonas pseudoalcaligenes
N_Geneseq_36:X20517	+	36.00	94.60	1.1e+03	6761	Polynucleotide sequence from t
N_Geneseq_36:T28774	+	36.00	88.22	2.6e+03	14042	Human placental calcium sens
N_Geneseq_36:V05995	+	36.00	88.22	2.6e+03	14042	Human placental calcium sens
N_Geneseq_36:T28776	+	36.00	88.21	2.6e+03	14044	Human parathyroid calcium sen
N_Geneseq_36:T05997	+	36.00	88.21	2.6e+03	14044	Human parathyroid calcium sen
N_Geneseq_36:T05997	+	36.00	88.19	2.6e+03	14080	Human kidney calcium sensor p
N_Geneseq_36:V05996	+	36.00	88.19	2.6e+03	14080	Human kidney calcium sensor p
N_Geneseq_36:T28773	+	36.00	88.19	2.6e+03	14086	Human calcium sensor protein
N_Geneseq_36:V05994	+	36.00	88.19	2.6e+03	14086	Human calcium sensor protein
N_Geneseq_36:T07264	+	35.00	108.70	187.15	900	Mutant LAG1 coding sequence. Eu
N_Geneseq_36:T43690	+	35.00	103.65	357.68	1604	Alcohol acetyltransferase cod
N_Geneseq_36:T15232	+	35.00	103.18	380.25	1694	Tumor necrosis factor recepto
N_Geneseq_36:T94635	+	35.00	103.18	380.25	1694	TNF-RI-DP ligand protein clone
N_Geneseq_36:T07263	+	35.00	101.16	492.57	2134	LAG1 coding sequence. Eukaryot
N_Geneseq_36:T94639	+	35.00	99.87	581.08	2473	TNF-RI-DP ligand protein clone
N_Geneseq_36:V23687	+	35.00	97.05	834.34	3415	L. lactis DB1341 pfl gene. Lac
N_Geneseq_36:Q62924	+	35.00	88.66	2.4e+03	8920	Carbamoyl-phosphate-synthetase
N_Geneseq_36:X20546	+	35.00	83.21	4.9e+03	16636	Polynucleotide sequence from
N_Geneseq_36:X20569	+	35.00	81.31	6.3e+03	20682	Polynucleotide sequence from
N_Geneseq_36:T80649	+	34.00	118.82	51.14	189	Type II topoisomerase database
N_Geneseq_36:X30892	+	34.00	108.83	184.22	593	Streptococcus pneumoniae genom
N_Geneseq_36:T96357	+	34.00	107.89	207.71	660	CD28 cDNA. New xanthene derivat

N_Geneseq_36:X20213 - 34.00 106.68 242.58 758 ! Enterococcus faecalis EF110
N_Geneseq_36:Q28837 - 34.00 106.63 244.01 762 ! Sequence of the CD28 gene. S
N_Geneseq_36:X19109 + 34.00 106.20 258.05 801 ! Rhodospiridium toruloides D-
N_Geneseq_36:X20212 - 34.00 105.30 289.67 888 ! Enterococcus faecalis gene E

seq_name: N_Geneseq_36:T84234

seq_documentation_block:

```
ID T84234 standard; DNA; 3110 BP.  
AC T84234;  
DT 01-SEP-1998 (first entry)  
DE DNA encoding an autolysin and 3 unknown proteins  
KW Staphylococcus aureus protein; ribozyme; antiseptic sequence; control;  
KW Staphylococcus aureus; regulatory element; bacterial gene expression;  
KW vaccine; Staphylococcus aureus; food poisoning; scaled skin syndrome;  
KW toxic shock syndrome; ss.  
OS Staphylococcus aureus.  
FH Key Location/Qualifiers  
FT CDS 579..1272  
FT CDS /tag= a  
FT CDS 1668..1844  
FT CDS /tag= b  
FT CDS 1885..2064  
FT CDS /tag= c  
FT CDS 2326..2832  
FT CDS /tag= d  
FT CDS /product= "autolysin"
```

WO9730070-A1.

21-AUG-1997.

19-FEB-1997; U023118.

20-FEB-1996; US-011888.

(SMIK) SMITHKLINE BEECHAM CORP.

Black Mt, Burnham MK, Hodgson JB, Knowles DJC, Nicholas RO,

Pratt JM, Reichard RW, Rosenberg M, Ward JM;

WPI; 97-424969/39.

P-PSDB: W28340, W28341, W28342, W28343.

Novel polypeptide(s) from Staphylococcus aureus strain WCUH29 - used

to isolate antimicrobial compounds, and in vaccines against S.

aureus infection

Claim 9; Page 977-978; 989pp; English.

The present sequence encodes 3 Staphylococcus aureus proteins of

unknown function and a protein, that, based on homology with a

known Staphylococcus aureus protein, is believed to be an autolysin

(EC 3.5.1.28) (N-acetylmuramoyl-L-alanine amidase).

The present sequence was obtained from a

library of clones of S. aureus WCUH 29 in Escherichia coli. The DNA

sequence can be used in the construction of Staphylococcus aureus

sequences to control the expression of Staphylococcus aureus

sequence is also useful as a source of regulatory elements for the

control of bacterial gene expression. The encoded protein may be used

to produce vaccines to enable a host to produce specific antibodies

with antibacterial action. These vaccines and antibodies would protect

a host against invasion by S. aureus, and conditions relating to

Staphylococcus aureus infection, e.g. Staphylococcus aureus food poisoning, scaled

skin syndrome, and toxic shock syndrome.

Sequence 3110 BP; 1111 A; 488 C; 675 G; 833 T;

alignment_scores:

Quality: 40.00 Length: 14

Ratio: 3.333 Gaps: 0

Percent Similarity: 85.714 Percent Identity: 35.714

alignment_block:

US-08-653-294-20 x T84234 ..

Align seg 1/1 to: T84234 from: 1 to: 3110

1 TyrArgLeuAlaIleArgIle***ArgIleLeuArgTyr 14

|||||:|||||:|||||:|||||:|||||:|||||

785 TACCAGGTAGGTTGAAGTATGATGAGGTTATTCGGTAT 826

seq_name: N_Geneseq_36:V53557

```

seq_documentation_block:
ID V53557 standard; DNA; 3110 BP.
AC V53557.
DT 30-OCT-1998 (first entry)
DE DNA encoding 2 Staphylococcus aureus proteins of unknown function.
KW Staphylococcus aureus protein; immune response induction; eye infection;
KW antibody production; T-cell immune response; gastrointestinal infection;
KW respiratory infection; inhibitor; bacterial infection; cardiac infection;
KW central nervous system; kidney infection; urinary tract infection;
KW antimicrobial compound identification; broad spectrum antibiotic;
KW therapy; ss.
OS Staphylococcus aureus.
FH Key Location/Qualifiers
FT 1208..1300
CDS
FT /*tag= a
FT /note= "encoded protein shown in W77771"
FT 1282..1668
FT /*tag= b
FT /note= "encoded protein shown in W77772"
PN EP-841394-A2.
PD 13-MAY-1998.
PE 24-SEP-1987; 307485.
PR 24-SEP-1996; US-027032.
PA (SMIK ) SMITHKLINE BEECHAM CORP.
PI Black MT, Burnham MKR, Hodgson JE, Knowles DJC,
PI Lonetto MA, Nicholas RO, Pratt JM, Reichard RW, Rosenberg M,
PI Ward JM,
PI WPI: 98-252940/23.
DR P-PSDB; W7771, W7772.
PT New nucleic acid sequences from Staphylococcus aureus WCHU29 -
PT useful in vaccines and for treatment of bacterial infections of e.g.
PT respiratory tract and central nervous system
PS Claim 1: Page 214-216; 390pp; English.
CC This sequence encodes 2 Staphylococcus aureus proteins of unknown
CC function, and represents a DNA sequence of the invention.
CC The DNA sequences were isolated from Staphylococcus aureus WCHU29
CC (NCIMB 40771). Host cells containing the DNA sequences are used to
CC produce polypeptides or fragments. The proteins are used in the treatment
CC of disease, for inducing an immune response by administering them, to
CC produce antibody and/or T-cell immune response. Antagonists of the
CC proteins are used for the inhibition of bacterial polypeptides.
CC Conditions which may be treated include bacterial infections, especially
CC respiratory, cardiac, gastrointestinal, central nervous, eye, kidney,
CC urinary tract, skin, bones and joints. The proteins can also be used to
CC identify antimicrobial compounds which are broad spectrum antibiotics,
CC especially useful in the treatment of H. Pylori infection.
SQ Sequence 3110 BP; 1111 A; 488 C; 576 G; 832 T;

alignment_scores:
Quality: 40.00 Length: 14
Ratio: 3.33 Gaps: 0
Percent Similarity: 85.714 Percent Identity: 35.714

alignment_block:
US-08-653-294-20 x V53557 ..
Align seg 1/1 to: V53557 from: 1 to: 3110
1 TyrArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
|||||:|||||:|||||: |||:|||||
785 TACCAGGTACGTTGAAGTATGAGGTTATTTATCGGTAT 826

seq_name: N_Geneseq_36:V74331
seq_documentation_block:
ID V74331 standard; DNA; 10470 BP.
AC V74331.
DT 16-MAR-1999 (first entry)
DE Staphylococcus aureus contig SEQ ID #20.
KW Computer readable medium; vaccine; S.aureus infection; Immunodetection;

```

cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;
skin infection; surgical wound infection; scalded skin syndrome;
toxic shock syndrome; ds.
Staphylococcus aureus.
Key Location/Qualifiers
misc_feature 361..420
/*tag= a
/note= "these bases represent a line of missing text in
the sequence listing in the specification. They
are included to maintain the nucleotide numbering
given in the specification for this DNA sequence"
misc_feature 2161..2220
/*tag= b
/note= "these bases represent a line of missing text in
the sequence listing in the specification. They
are included to maintain the nucleotide numbering
given in the specification for this DNA sequence"
misc_feature 3961..4020
/*tag= c
/note= "these bases represent a line of missing text in
the sequence listing in the specification. They
are included to maintain the nucleotide numbering
given in the specification for this DNA sequence"
misc_feature 5761..5820
/*tag= d
/note= "these bases represent a line of missing text in
the sequence listing in the specification. They
are included to maintain the nucleotide numbering
given in the specification for this DNA sequence"
misc_feature 7561..7620
/*tag= e
/note= "these bases represent a line of missing text in
the sequence listing in the specification. They
are included to maintain the nucleotide numbering
given in the specification for this DNA sequence"
misc_feature 9361..9420
/*tag= f
/note= "these bases represent a line of missing text in
the sequence listing in the specification. They
are included to maintain the nucleotide numbering
given in the specification for this DNA sequence"
EP-786519-A2.
30-JUL-1997.
07-JAN-1997; 100117.
05-JAN-1996; US-009861.
(HUMA-) HUMAN GENOME SCI INC.
Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA,
Rosen CA,
WPI: 97-374922/35.
Polynucleotide(s) and proteins derived from Staphylococcus aureus -
stored on computer readable medium and used in the production of
anti-S.aureus vaccines
Claim 1: Page 271-277; 3271pp; English.
This sequence represents one of 5191 Staphylococcus aureus DNA sequences
of the invention. The DNA sequences are recorded on a computer readable
medium, preferably selected from a floppy or hard disk, random access
memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
the S.aureus DNA sequences allows putative functions to be assigned so
that protein-encoding or regulatory regions of commercial, therapeutic or
industrial importance can be obtained. Specifically, sequences which are
likely to encode antigens have been identified and these polypeptides can
be used in a vaccine composition against S.aureus infection. The
polypeptides can also be used in a kit for the immunodetection of
S.aureus in a sample. S.aureus is implicated in numerous human diseases,
including cellulitis, eyelid infections, food poisoning, osteomyelitis,
skin and surgical wound infections, scalded skin syndrome, toxic shock
syndrome, etc. Organisms transformed with the DNA sequences can be used
for recombinant production of the polypeptides. The new DNA sequences
(and their fragments) are useful as primers or probes for isolating
homologues of any of the S.aureus DNA sequences contained on the
computer readable medium.
Sequence 10470 BP; 2804 A; 2009 C; 1550 G; 3742 T;

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alignment_scores:
  Quality: 40.00      Length: 14
  Ratio: 3.333       Gaps: 0
  Percent Similarity: 85.714   Percent Identity: 35.714

alignment_block:
US-08-653-294-20 x V74331/rev ..
  Align seg 1/1 to reverse of: V74331 from: 1 to: 10470
    1 TyrArgLeuAlaIleArgIle***ArgIleLeuArgTyr 14
      |||:|||||:|||||: |||:|||||: |||:|||||
    9287 TACAGGTAGCTTGAAGATGATGAGGTTATTATCGGTAT 9246

seq_name: N_Geneseq_36.V69717

seq_documentation_block:
ID V69717 standard; cDNA: 4031 BP.
AC V69717;
DT 01-MAR-1999 (first entry)
DE Tumour rejection antigen precursor MAGE-C1 cDNA.
KW MAGE-C1; human; tumour rejection antigen precursor; TRAP;
KW therapy; diagnosis; ds.
OS Homo sapiens.
FH Key
FT CDS
FT FT
FT FT
PN WO9849184-A1.
PD 05-NOV-1998;
PF 24-APR-1998; U08493.
PR 25-APR-1997; US-845528.
PA (LUDW-) LUDWIG INST CANCER RES.
PI Boon-Falleur T, De Smet C, Lucas S;
DR WPI: 99-024041/02.
PT Tumour rejection antigen precursors - used for determining presence
PT of cytolytic T cells specific for complexes of a human leukocyte
PT antigen
PS Claim 26; Page 40-42; 84pp; English.
CC This nucleotide sequence comprises novel human tumour rejection
CC antigen precursor (TRAP) MAGE-C1 cDNA (see also V69720). MAGE-C1
CC is a novel member of the MAGE family that may be recognised by
CC cytotoxic T cells, leading to lysis of the tumour cells which
CC express it. MAGE-C1 and MAGE-C2 (see W81546-47) are expressed in a
CC variety of tumours and in normal testis cells, but not by other
CC normal cells. The MAGE-C1 cDNA was isolated from a melanoma
CC LB373-MEL cDNA library using a probe generated from LB-373-MEL
CC RNA by PCR (see V69728-29). It shows homology to MAGE-A1 cDNA (see
CC V69719) and codes for a putative 1072-amino acid protein. The
CC MAGE-C1 gene was localised to chromosome region Xq26-q27. MAGE-C1
CC and MAGE-C2 cDNAs (see V69726) are claimed, as are: expression
CC vectors; transformed or transfected cell lines (e.g. COS and CHO);
CC an isolated TRAP encoded by the cDNAs; a kit useful in a PCR based
CC assay comprising an oligonucleotide having a sequence of
CC nucleotides 18-34 of the 4031 bp MAGE-C1 cDNA and an oligonucleotide
CC having a sequence which is complementary to nucleotides 200-217 of
CC the 4031 bp cDNA sequence; a method for determining expression of
CC a MAGE-C1 gene using the kit; a polypeptide comprising a number of
CC tumour rejection antigens derived from MAGE-C1 or MAGE-C2; and a
CC polypeptide comprising at least one tumour rejection antigen derived
CC from MAGE-C1 or MAGE-C2 and at least one other tumour rejection
CC antigen. MAGE-C1 and MAGE-C2 can be used in a method for determining
CC the presence of cytolytic T cells specific for complexes of a human
CC leukocyte antigen (HLA).
SQ Sequence 4031 BP; 849 A; 1138 C; 877 G; 1167 T;

alignment_scores:
  Quality: 39.00      Length: 13
  Ratio: 3.545       Gaps: 0
  Percent Similarity: 84.615   Percent Identity: 46.154

alignment_block:
US-08-653-294-20 x V69720 ..
  Align seg 1/1 to: V69720 from: 1 to: 4225
    1 TyrArgLeuAlaIleArgIle***ArgIleLeuArg 13
      |||:|||||:|||||: |||:|||||: |||:|||||
    1577 TACAGGTTCCTCGAGAGTCTCAAGAGTCTTTTGGAG 1565

seq_name: N_Geneseq_36.T48660

seq_documentation_block:

```

```

US-08-653-294-20 x V69717 ..
  Align seg 1/1 to: V69717 from: 1 to: 4031
    1 TyrArgLeuAlaIleArgIle***ArgIleLeuArg 13
      |||:|||||:|||||: |||:|||||: |||:|||||
    1317 TACAGAGTTCCTCGAGAGTCTCAAGAGTCTTTTGGAG 1355

seq_name: N_Geneseq_36.V69720

seq_documentation_block:
ID V69720 standard; cDNA: 4225 BP.
AC V69720;
DT 01-MAR-1999 (first entry)
DE Tumour rejection antigen precursor MAGE-C1 cDNA.
KW MAGE-C1; human; tumour rejection antigen precursor; TRAP;
KW therapy; diagnosis; ds.
OS Homo sapiens.
FH Key
FT CDS
FT FT
FT FT
PN WO9849184-A1.
PD 05-NOV-1998;
PF 24-APR-1998; U08493.
PR 25-APR-1997; US-845528.
PA (LUDW-) LUDWIG INST CANCER RES.
PI Boon-Falleur T, De Smet C, Lucas S;
DR WPI: 99-024041/02.
PT Tumour rejection antigen precursors - used for determining presence
PT of cytolytic T cells specific for complexes of a human leukocyte
PT antigen
PS Claim 1; Page 48-50; 84pp; English.
CC This nucleotide sequence encodes novel human tumour rejection
CC antigen precursor (TRAP) MAGE-C1 (see W81546). MAGE-C1 is a novel
CC member of the MAGE family that may be recognised by cytotoxic T
CC cells, leading to lysis of the tumour cells which express it.
CC MAGE-C1 and MAGE-C2 (see W81547) are expressed in a variety of
CC tumours and in normal testis cells, but not by other normal cells.
CC The MAGE-C1 cDNA was isolated from a melanoma LB373-MEL cDNA by
CC PCR amplification (see V69732-33). It shows homology to MAGE-A1
CC cDNA (see V69719). The MAGE-C1 gene was localised to Xq26-q27.
CC MAGE-C1 and MAGE-C2 cDNAs (see V69726) are claimed, as are:
CC expression vectors; transformed or transfected cell lines (e.g. COS
CC and CHO); an isolated TRAP encoded by the cDNAs; a kit useful in a
CC PCR based assay; a method for determining expression of a MAGE-C1
CC gene using the kit; a polypeptide comprising a number of tumour
CC rejection antigens derived from MAGE-C1 or MAGE-C2; and a polypeptide
CC comprising at least one tumour rejection antigen derived from
CC MAGE-C1 or MAGE-C2 and at least one other tumour rejection antigen.
CC MAGE-C1 and MAGE-C2 can be used in a method for determining the
CC presence of cytolytic T cells specific for complexes of a human
CC leukocyte antigen (HLA).
SQ Sequence 4225 BP; 871 A; 1198 C; 923 G; 1233 T;

alignment_scores:
  Quality: 39.00      Length: 13
  Ratio: 3.545       Gaps: 0
  Percent Similarity: 84.615   Percent Identity: 46.154

alignment_block:
US-08-653-294-20 x V69720 ..
  Align seg 1/1 to: V69720 from: 1 to: 4225
    1 TyrArgLeuAlaIleArgIle***ArgIleLeuArg 13
      |||:|||||:|||||: |||:|||||: |||:|||||
    1577 TACAGAGTTCCTCGAGAGTCTCAAGAGTCTTTTGGAG 1565

seq_name: N_Geneseq_36.T48660

seq_documentation_block:

```



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PA (KAOS ) KAO CORP.
PI Ara K, Hatada Y, Ito S, Kawai S, Ozaki K;
DR WPI: 97-118708/11.
DR P-PSDB: W11326.
PT DNA encoding alkaline liquefying alpha-amylase - useful in
PT dish-washing and laundry detergents for removal of starch dirt
PS Claim 8; Page 23-26; 40pp; English.
CC This sequence represents the coding sequence for an alkaline liquefying
CC alpha-amylase. Alpha-amylase is an enzyme that acts on starch-related
CC polysaccharides, hydrolysing the alpha-1,4-glucoside bond of the
CC polysaccharide molecule. Alkaline liquefying alpha-amylases exhibit
CC resistance to surfactants used in detergents, and decompose starch or
CC starch-related polysaccharides in a highly random manner. The Bacillus
CC species KSM-Ap1378, from which this sequence was isolated, is an
CC alkalophilic Bacillus strain. It was isolated from soil in the vicinity
CC of the city of Tohigi. The enzyme is useful in improving the efficiency
CC of dish-washing and laundry detergents, particularly on starch dirt.
SQ Sequence 1776 BP; 575 A; 305 C; 417 G; 479 T;

alignment_scores:
Quality: 37.00 Length: 13
Ratio: 3.700 Gaps: 0
Percent Similarity: 76.923 Percent Identity: 69.231

alignment_block:
US-08-653-294-20 x T51339 ..
Align seg 1/1 to: T51339 from: 1 to: 1776

2 ArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
||||| :||||| ||| |||||||||
1320 AGGTTACCTTCGGTATTTACGGTACTTACGGTAT 1358

seq_name: N_Geneseq_36:N60309
seq_documentation_block:
ID N60309 standard; DNA; 2109 BP.
AC N60309;
DT 10-JUN-1991 (first entry)
DE Equine IFN-omega-1 from PAH61.
KW IFN-omega-1; equine; Interferon; ss.
OS Equus caballus.
FH Key Location/Qualifiers
FT cds /*tag= a
FT signal_peptide /*product= IFN-omega-1
FT mat_peptide /*tag= b
FT /*tag= c
EP-186098-A.
PD 02-JUL-1986.
PF 17-DEC-1985; 116083.
PR 18-DEC-1984; DE-446122.
PA (BOEH ) BOEHRINGER INGELHEIM.
PI Himmler A, Hauptmann R, Haeufel N, Adolf G, Swetley P;
DR WPI: 86-170649/27.
DR P-PSDB: P60400.
PT New equine and canine interferon - and recombinant DNA molecules
PT coding for them, and transformed cells.
PS Disclosure; Fig 12; 149pp; German.
CC Microorganisms transformed with the recombinant sequence produce IFN
CC which is useful therapeutically in veterinary medicine.
CC See also N60306-13 and N60938.
SQ Sequence 2109 BP; 609 A; 476 C; 471 G; 553 T;

alignment_scores:
Quality: 37.00 Length: 14
Ratio: 3.364 Gaps: 0
Percent Similarity: 78.571 Percent Identity: 50.000

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alignment_block:
US-08-653-294-20 x N60309/rev ..
Align seg 1/1 to reverse of: N60309 from: 1 to: 2109

1 TyrArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
||||| :||||| ||| |||||||||
1896 TATTTCTTCTCTGTCAGGTACAGCGGATTCCTCGTAAGTAC 1855

seq_name: N_Geneseq_36:T33646
seq_documentation_block:
ID T33646 standard; DNA; 2555 BP.
AC T33646;
DT 11-DEC-1996 (first entry)
DE Aspergillus arabinofuranosidase gene.
KW Arabinofuranosidase; Abfc; arabinoxylan; viscosity modifier; food;
KW feedstuff; ds.
OS Aspergillus niger strain 3M43.
FH Key Location/Qualifiers
FT promoter 1..869
FT /*tag= a
FT cds 870..1757
FT /*tag= b
FT signal_peptide 870..947
FT /*tag= c
FT mat_peptide 948..1754
FT /*tag= d
FT terminator 1755..2555
FT /*tag= e
WO9629416-A1.
PN 26-SEP-1996.
PD 11-MAR-1996; E01009.
PR 17-MAR-1995; GB-005479.
PA (DANI-) DANISCO AS.
PI Baruch A, Madrid SM, Rasmussen P;
DR WPI: 96-443191/44.
DR P-PSDB: W00810.
PT Aspergillus arabinofuranosidase - useful for degradation of
PT arabinoxylan
PS Example; Page 52-55; 105pp; English.
CC The Aspergillus niger 3M43 arabinofuranosidase gene (T33646)
CC codes for the precursor form (W00810) of the arabinofuranosidase
CC enzyme. It was isolated from a genomic DNA library by screening
CC with a PCR clone obtd. by amplification of A. niger DNA using
CC primers (see also T33644-45) based on an isolated peptide (W00806)
CC the enzyme. The mature enzyme coding sequence (see also T33640)
CC can be used for prodn. of arabinofuranosidase (W04167) in
CC transformed hosts, pref. Aspergillus or a transgenic plant. The
CC promoter (see also T33641), terminator (T33642) and signal sequence
CC (T33643) may also be used to control expression of the
CC arabinofuranosidase gene, or other gene of interest, in a host
CC cell.
SQ Sequence 2555 BP; 632 A; 651 C; 615 G; 657 T;

alignment_scores:
Quality: 37.00 Length: 12
Ratio: 3.700 Gaps: 0
Percent Similarity: 83.333 Percent Identity: 58.333

alignment_block:
US-08-653-294-20 x T33646 ..
Align seg 1/1 to: T33646 from: 1 to: 2555

1 TyrArgLeuAlaIleArgIle***ArgIleLeuLeu 12
||||| :||||| ||| |||||||||
165 TACCAACTTTTGTTCGATTGATCAGATACATTG 200

seq_name: N_Geneseq_36:V29571
seq_documentation_block:

```

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ID V29571 standard; DNA; 4815 BP.
AC V29571;
DE 04-AUG-1998 (first entry)
DE L. lactis soluble part (F1) exhibiting ATPase activity encoding DNA.
KW Biomass production; uncoupled ATPase; F0F1 ATPase; membrane bound;
KW F1; Lactococcus lactis; ds.
OS Lactococcus lactis.
FH Key
FT Location/Qualifiers
FT 26..553
FT /tag= a
FT /product= "ATPase subunit"
FT /gene= "atpB"
FT /note= "delta subunit of the F1 portion of F0F1 ATPase"
FT 742..2244
FT /tag= b
FT /product= "ATPase subunit"
FT /gene= "atpA"
FT /note= "alpha subunit of the F1 portion of F0F1 ATPase"
FT 2260..3129
FT /tag= c
FT /product= "ATPase subunit"
FT /gene= "atpG"
FT /note= "gamma subunit of the F1 portion of F0F1 ATPase"
FT 3301..4710
FT /tag= d
FT /product= "ATPase subunit"
FT /gene= "atpD"
FT /note= "beta subunit of the F1 portion of F0F1 ATPase"
PN W09810089-A1.
PD 12-MAR-1998.
PF 08-SEP-1997; DK0373.
PA 06-SEP-1996; DK-000963.
PA (JENS/) JENSEN P R.
PI Snoep JL, Westerhoff HV;
DR WPI: 98-193637/17.
DR P-PSDB; W56790, W56791, W56792, W56793.
DR Method improving production of biomass or a desired product - by
DR expressing an uncoupled ATPase activity in the cell
PS Claim 16; Pages 35-41; 78pp; English.
CC This DNA encodes the soluble part (F1) of membrane bound (F0F1 type) H+
CC - ATPase or a portion of F1 exhibiting ATPase activity. The DNA is
CC derived from Lactococcus lactis subsp. cremoris strain MGL363. This is
CC used in a novel method for improving the production of biomass or a
CC desired product from a cell. The method comprises expressing an uncoupled
CC ATPase activity in the cell to induce conversion of ATP to ADP without
CC primary effects on other cellular metabolites or functions and incubating
CC the cell with a suitable substrate to produce the biomass or product. The
CC expression is directed using a vector including DNA encoding the soluble
CC part (F1) of the membrane bound (F0F1 type) H+-ATPase or a portion of F1
CC exhibiting ATPase activity, the DNA being derived from Lactococcus lactis
CC subsp. cremoris, Lactococcus lactis subsp. Lactis, Streptococcus
CC thermophilus, Phaffia rhodozyma or Trichoderma reesei, where the DNA is
CC under the control of a promoter. An ideal ATPase is the membrane bound
CC H+ ATPase. This enzyme complex consists of two parts, the membrane
CC integral part and the (F0) and the cytoplasmic part (F1). Together the
CC two parts couple the hydrolysis of ATP and ADP to the translocation of
CC protons across the cytoplasmic membrane, or vice versa. The proton
CC gradient is used to drive ATP synthesis from ADP and Pi. The method can
CC be used for optimising the formation of biomass or a desired product,
CC e.g. the product may be lactic acid which results in the acidification of
CC dairy products. The method is more efficient than currently used methods
CC of biomass production.
SQ Sequence 4815 BP; 1491 A; 897 C; 981 G; 1446 T;

alignment_scores:
Quality: 37.00 Length: 14
Ratio: 3.083 Gaps: 0
Percent Similarity: 85.714 Percent Identity: 42.857

alignment_block:
US-08-653-294-20 x V29571
..

Align seg 1/1 to: V29571 from: 1 to: 1446 T;

1 TyrArgLeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
4384 TATGAAGTTCAGTGAAGTTCACGTCGCTTCACAGCTAC 4425

seq_name: N_Geneseq_36:V52237

seq_documentation_block:
ID V52237 standard; DNA; 6735 BP.
AC V52237;
DE 23-OCT-1998 (first entry)
DE Streptococcus pneumoniae genome fragment SEQ ID NO:104.
KW Streptococcus pneumoniae; S. pneumoniae; genome; diagnosis; assay;
KW computer readable medium; vaccine; pharmaceutical composition; ds.
OS Streptococcus pneumoniae.
PN W09818931-A2.
PD 07-MAY-1998.
PF 30-OCT-1997; U19588.
PR 31-OCT-1996; US-029960.
PA (HUNA-) HUMAN GENOME SCI INC.
PI Barash SC, Choi GH, Dillon PJ, Dougherty BA, Fannon M,
PI Kunsch CA, Rosen CA;
DR WPI: 98-272225/24.
DR Computer-readable medium with recorded Streptococcus pneumoniae
DR polynucleotide sequences - useful in diagnostic kits and assays, and
DR pharmaceutical compositions and vaccines for Streptococcus
DR pneumoniae
PS Claim 1; Page 773-777; 1409pp; English.
CC The present invention describes a computer readable medium which has
CC the nucleotide sequences SEQ ID NO:1 to 391 (V52134 to V52524) recorded
CC on it, or a representative fragment or a sequence at least 95% identical
CC to SEQ ID NO: 1 to 391. The nucleotide sequences depicted in SEQ ID NO:1
CC to 391 (V52134 to V52524) are genomic fragments from Streptococcus
CC pneumoniae. The present invention also describes an isolated nucleic acid
CC molecule encoding a homologue of any of the fragments of the S.pneumoniae
CC genome (SEQ ID NO:1 to 391) where the nucleic acid molecule is produced
CC by a process comprising: (a) screening a genomic DNA library using as a
CC probe a target sequence defined by any of the sequences in SEQ ID NO:1
CC to 391, identifying members of the library which contain sequences
CC that hybridise to the target sequence and isolating the nucleic acid
CC molecules from the members; or (b) isolating mRNA, DNA or cDNA produced
CC from an organism, amplifying nucleic acid molecules whose nucleotide
CC sequence is homologous to amplification primers derived from the
CC fragment of the S. pneumoniae genome to prime the amplification and
CC isolating the amplified sequences. The computer readable medium can be
CC used in a computer-based system for identifying fragments of the
CC S. pneumoniae genome of commercial importance, or expression modulating
CC fragments of the S. pneumoniae genome. Products from the present
CC invention can be used in diagnosis kits and assays, and pharmaceutical
CC compositions and vaccines for S. pneumoniae.
SQ Sequence 6735 BP; 2122 A; 1141 C; 1415 G; 2057 T;

alignment_scores:
Quality: 37.00 Length: 12
Ratio: 3.700 Gaps: 0
Percent Similarity: 83.333 Percent Identity: 66.667

alignment_block:
US-08-653-294-20 x V52237
..

Align seg 1/1 to: V52237 from: 1 to: 6735

3 LeuAlaIleArgIle***ArgIleLeuLeuArgTyr 14
|||||:|||||:|||||:|||||:|||||:|||||:|||||:
5344 TTATCAATAAATCCAGAGATATGCTATCTTAC 5379
```

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 01:29:43 ; Search time 133.56 seconds
(without alignments)
2.660 Million cell updates/sec

Title: US-08-653-294-21

Perfect score: 70

Sequence: 1 AYRLIKVIRVLKY 15

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 188963 seqs, 23686106 residues

Total number of hits satisfying chosen parameters: 188963

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database : A_Geneseq_36.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	37	52.9	497	1 W49912	Mouse CLK serine/threonine kinase
2	36	51.4	141	1 W04183	Cellubrevin-4, iso
3	36	51.4	1226	1 W13825	Yeast transcriptio
4	35	50.0	302	1 W76777	B. malayi ankryrin
5	35	50.0	302	1 W70609	Ankryrin protein PB
6	35	50.0	1784	1 R77223	Tuberous sclerosis
7	34	48.6	33	1 W76789	D. immitis pDlank1
8	34	48.6	37	1 W77378	Lytic peptide with
9	34	48.6	303	1 W76774	D. immitis ankryrin
10	34	48.6	303	1 W70606	Ankryrin protein fr
11	34	48.6	395	1 W82822	Pseudomonas mendoc
12	34	48.6	1594	1 P81184	Sequence encoded b
13	34	48.6	1745	1 W76776	D. immitis ankryrin
14	34	48.6	1745	1 W70608	Full length ankryri
15	33	47.1	20	1 R92907	HLA-B2702 CTL modu
16	33	47.1	20	1 R93428	HLA-B2702 84-75-84
17	33	47.1	20	1 W33778	Immunomodulating d
18	33	47.1	550	1 W81351	Human guanine nucl
19	33	47.1	580	1 W81349	Human guanine isom
20	33	47.1	878	1 Y00868	S. tuberosum isom
21	33	47.1	2861	1 W27227	Human TRIO phospho
22	32	45.7	12	1 R93429	HLA-B2702 84-79-84
23	32	45.7	12	1 W33798	Peptide B2702.84-7
24	32	45.7	12	1 W33799	Immunomodulating d
25	32	45.7	35	1 P90056	Human derived pept
26	32	45.7	35	1 R15605	ASP-5 analogue (3)
27	32	45.7	227	1 W22173	S. thermophilus exo
28	32	45.7	227	1 W14074	S. thermophilus exo
29	32	45.7	283	1 W41497	Avian cyclin C pro
30	32	45.7	325	1 W41496	Human cyclin C pro
31	32	45.7	374	1 W63694	Human secreted pro
32	32	45.7	443	1 W84264	Protein encoded by
33	32	45.7	626	1 W41501	Human DP.75, a put
34	32	45.7	2410	1 W19723	Cell cycle checkpo

35 32 45.7 2480 1 W19724 Cell cycle checkpo
36 32 45.7 2644 1 W13152 Human ataxia and r
37 32 45.7 2644 1 W84271 A human ATR protei
38 32 45.7 4472 1 R97246 Virulence gene clu
39 31.5 45.0 57 1 Y02689 Human secreted pro
40 31 44.3 23 1 P81601 Sequence of artifi
41 31 44.3 37 1 W83937 Human secreted pro
42 31 44.3 62 1 P80591 Sequence of artifi
43 31 44.3 101 1 R06693 Feline infectious
44 31 44.3 130 1 W90020 Expressed antigen
45 31 44.3 155 1 W98359 H. pylori GHPO 127

ALIGNMENTS

RESULT 1
ID W49912 standard; Protein: 497 AA.
AC W49912;
DT 20-JUL-1998 (first entry)
DE Mouse CLK serine/threonine kinase mCLK2.
KW mCLK2; CLK; serine/threonine kinase; protein kinase; LAMMER kinase;
KW signal transduction; cancer; contraceptive; mouse; therapy;
KW diagnosis.
OS Mus musculus.
FH Key Location/Qualifiers
FT Domain 28..45
FT Domain /note= "nuclear localisation domain"
FT Domain 163..479
FT Peptide /note= "catalytic domain"
FT Peptide 388..393
FT Peptide /note= "LAMMER motif"
PN W09748723-A2.
PD 24-DEC-1997.
PF 17-JUN-1997; IB0946.
PR 19-DEC-1996; US-034286.
PR 17-JUN-1996; US-019629.
PR 09-AUG-1996; US-023485.
PR 13-NOV-1996; US-030860.
PR 15-NOV-1996; US-030964.
PR (PLAC) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.
PR Aoki N, Chen Z, Kharitonkov AI, Kim YW, Nayler O,
PI Ullrich A, Wang HY;
PI WPI; 98-120302/11.
PT New phosphatase and kinase enzyme(s) - useful in the diagnosis and
PT treatment of signal transduction disorders
PT Claim 11; Fig 4; 138pp; English.
CC This polypeptide comprises novel mouse CLK serine/threonine kinase
CC mCLK2, from the CLK serine/threonine kinase family of proteins that
CC regulate RNA splicing in cells. mCLK2 cDNA was cloned from a mouse
CC embryo 11.5 p.c. 12AP cDNA library. The invention discloses the
CC discovery of novel protein kinases mCLK2, mCLK3 (see W49912) and
CC mCLK4 (see W49914) of mol.wt. 59.9, 58.5 and 57.2 kDa
CC respectively, as well as other novel proteins (see W49906-10)
CC involved in cellular signal transduction, and provides vectors,
CC host cells, purified recombinant proteins, methods for identifying
CC compounds that activate or inhibit the novel proteins, as well as
CC methods for the diagnosis and treatment of diseases associated with
CC the novel proteins. Overexpression of CLK serine/threonine kinases
CC has been implicated in certain types of cancer. Compounds that
CC inhibit their catalytic activity or disrupt their interactions
CC with natural binding partners may act as anti-cancer therapeutics.
CC mCLK related molecules and compounds may also be useful as male
CC contraceptives.
SQ Sequence 497 AA;

Query Match 52.9%; Score 37; DB 1; Length 497;

Best Local Similarity 53.8%; Pred No. 29;

Matches 7; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 3 RLKIKVIRVLKY 15

Db 188 RVLVKIRNVEKY 200
|:|:|:|:|:|:|

RESULT 2

W04183
ID W04183 standard; Protein; 141 AA.
AC W04183;
DT 29-MAY-1997 (first entry)
DE Cellubrevin-4.
KW CB-1; CB-2; CB-3; CB-4; human; cellubrevin; study; regulation;
KW vesicle trafficking; diagnosis; treatment; infection; leukaemia;
KW traumatic tissue damage; asthma; arthritis; cancer; lymphoma.
OS Homo sapiens.
PN W09629407-A2.
PD 26-SEP-1996.
PF 22-MAR-1996; U03835.
PR 23-MAR-1995; US-409373.
PA (INCY-) INCYTE PHARM INC.
PI Hawkins PR, Murry LE, Seilhamer JJ, Stuart SG;
DR WPI: 96-443183/44.
DR N-PSDB: T33717.
PT Isolated human cellubrevin polynucleotide(s) - useful to develop
PT prods. for diagnosis and treatment of conditions involving abnormal
PT membrane trafficking
PS Claim 25; Page 32; 59pp; English.
CC W04180-83 are novel human cellubrevins (CB-1, -2, -3 and -4). CBS
CC can be used for the study and regulation of vesicle trafficking in
CC normal, and in acute and chronic disease situations, and for the
CC diagnosis and treatment of conditions caused by infection, traumatic
CC tissue damage, hereditary disease, e.g. asthma or arthritis, invasive
CC cancer, leukaemia and lymphoma or other physiologic/pathologic problems
CC associated with induced, and otherwise abnormal, membrane trafficking.
CC In particular, the CB-4 polynucleotide (isolated from a cerebellum
CC library (CBUN0701) can be used in a diagnostic test for conditions or
CC diseases in which its expression is induced, e.g. cerebellar degenerative
CC diseases or brain tumours, while CB-4 inhibitors can be used to treat
CC such conditions or diseases
SQ Sequence 141 AA;

Query Match 51.4%; Score 36; DB 1; Length 141;
Best Local Similarity 53.3%; Pred. No. 12;
Matches 8; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

OY 1 AYRLIKVIRVLKY 15
|:|:|:|:|:|:|

Db 125 AILLVILIVMKY 139

RESULT 3

W13825
ID W13825 standard; Protein; 1226 AA.
AC W13825;
DT 04-JUN-1997 (first entry)
DE Yeast transcription regulatory factor SRB8.
KW Transcription regulatory factor; suppressor of RNA polymerase B;
KW SRB8; RNA polymerase II; holoenzyme; SWI/SNF.
OS Saccharomyces cerevisiae.
PN W09708301-A1.
PD 06-MAR-1997.
PF 28-AUG-1996; U14192.
PR 31-AUG-1995; US-521872.
PR 11-OCT-1995; US-540804.
PR 26-JAN-1996; US-590399.
PA (WHED) WHITEHEAD INST BIOMEDICAL RES.
PI Chao DM, Koleske AJ, Thompson CM, Young RA;
DR WPI: 97-179258/16.
DR N-PSDB: T59908.
PT Purified RNA polymerase II holoenzyme - comprises RNA polymerase II
PT and one or more regulatory proteins, pref. suppressor of RNA
PT polymerase B proteins or SWI/SNF proteins
PS Claim 11; Fig 10a-b; 154pp; English.

CC Novel yeast SRB (suppressor of RNA polymerase B) proteins SRB4, SRB5,
CC SRB6, SRB7, SRB8, SRB9, SRB10 and SRB11 (W13821-28) are transcription
CC regulatory factors that act as positive and negative regulators of
CC RNA polymerase II activity, and are components of the RNA polymerase
CC II holoenzyme. They were identified using methods designed to
CC identify transcription factors involved in RNA polymerase II
CC C-terminal domain (CTD) function. SRB8 and SRB9 appear to repress
CC CTD activity. Genomic clones (T59904-11) for the SRBs have been obtd.
CC SRBs can be used to treat diseases resulting from alteration or
CC deletion of the SRB gene, pref. by gene transfer technology. They
CC can also be used in in vitro transcription of DNA and to identify
CC cpds. that modify gene transcription.
SQ Sequence 1226 AA;

Query Match 51.4%; Score 36; DB 1; Length 1226;
Best Local Similarity 77.8%; Pred. No. 1.1e+02;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 2 YRLIKVIR 10
|:|:|:|:|

Db 999 YHLLIKIIR 1007

RESULT 4

W76777
ID W76777 standard; Protein; 302 AA.
AC W76777;
DT 15-JAN-1999 (first entry)
DE B. malayi ankyrin pBmank302 protein.
KW Ankyrin; helminth; parasite; vaccine; infection;
KW passive immunogen; cytotoxic agent.
OS Dirofilaria immitis.
PN US5827692-A.
PD 27-OCT-1998.
PF 24-APR-1997; 847429.
PR 24-APR-1997; US-847429.
PA (HESK-) HESKA CORP.
PI Blehm ES, Tang L;
DR WPI: 98-593992/50.
DR N-PSDB: V63024.
PT Nucleic acids encoding ankyrins from helminth parasites - useful for
PT recombinant production of the proteins for use as vaccines and
PT treatments against helminth infection
PS Claim 8; Column 137-140; 84pp; English.
CC W76769-W76777 represent ankyrin proteins isolated from the helminth
CC parasites Dirofilaria immitis and Brugia malayi. The nucleic acids and
CC recombinant products are useful for the recombinant production of the
CC ankyrin polypeptides. These proteins can then be used as vaccines against
CC parasitic helminth, e.g. D. immitis or B. malayi. They can also be used
CC for therapy after infection, and to raise antibodies, also for use in
CC therapeutics, as passive immunogens, or as therapeutics against helminths
CC on conjugation to cytotoxic agents. The nucleic acids contained in
CC viruses, may also be used as viral vaccines, and the nucleic acids
CC themselves or in vectors may be used as genetic vaccines.
SQ Sequence 302 AA;

Query Match 50.0%; Score 35; DB 1; Length 302;
Best Local Similarity 60.0%; Pred. No. 40;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

OY 6 IKVIRIVLKY 15
|:|:|:|:|

Db 53 IKVVELLLKY 62

RESULT 5

W70609
ID W70609 standard; Protein; 302 AA.
AC W70609;
DT 21-JAN-1999 (first entry)
DE Ankyrin protein pBmank302.

KW Ankyrin protein; parasitic helminth; heartworm disease;
 KW elephantiasis; hydrocele.
 OS Brugia malayi.
 PN US5824306-A.
 PD 20-OCT-1998.
 PF 26-FEB-1998; 031485.
 PR 24-APR-1997; US-847429.
 PR 26-FEB-1998; US-031485.
 PA (HESK-) HESKA CORP.
 PI Blehm ES, Tang L;
 DR WPI: 98-593373/50.
 DR N-PSDB; V63315.
 FT Dirofilaria and Brugia ankyrin proteins and antibodies - useful for
 FT protection of animals from disease caused by parasitic helminth
 PS Claim 5; Columns 137-140; 84pp; English.
 CC The present sequence represents a Brugia malayi ankyrin protein.
 CC The ankyrin protein, or anti-ankyrin antibodies, may be used to
 CC protect an animal from disease caused by a parasitic helminth,
 CC especially where the disease is heartworm disease, elephantiasis or
 CC hydrocele.
 SQ Sequence 302 AA;

Query Match 50.0%; Score 35; DB 1; Length 302;
 Best Local Similarity 50.0%; Pred. No. 40;
 Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15
 III:IIII
 Db 53 IKVELLKY 62

RESULT 6
 R77223
 ID R77223 standard; Protein; 1784 AA.
 AC R77223.
 DT 04-DEC-1995 (first entry)
 DE Tuberosus sclerosis 2 TSC2 gene product.
 KW Tuberosus sclerosis 2; TSC2 gene; gene therapy; tumor.
 OS Homo sapiens.
 FH Key Location/Qualifiers
 FT modified_site 70
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT peptide 76..84
 FT /label= Repeat_motif
 FT peptide 99..107
 FT /label= Repeat_motif
 FT peptide 81..102
 FT /label= Leucine_zipper
 FT region 171..187
 FT /label= Membrane-spanning_region
 FT modified_site 132
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 154
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 211
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 311
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT modified_site 389
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"

FT modified_site 390
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 428
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT region 459..476
 FT /label= Membrane-spanning_region
 FT modified_site 526
 FT /label= Phosphorylation
 FT /note= "protein-kinase C phosphorylation
 FT site"
 FT region 555..572
 FT /label= Membrane-spanning_region
 FT modified_site 556
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 609
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT modified_site 655
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT modified_site 660
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase phosphorylation
 FT site"
 FT modified_site 716
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT modified_site 719
 FT /label= Phosphorylation
 FT /note= "potential tyrosine-kinase phosphorylation
 FT site"
 FT modified_site 757
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 790
 FT /label= Phosphorylation
 FT /note= "potential tyrosine-kinase phosphorylation
 FT site"
 FT region 804..821
 FT /label= Membrane-spanning_region
 FT modified_site 923
 FT /label= Phosphorylation
 FT /note= "potential protein-kinase C phosphorylation
 FT site"
 FT modified_site 932
 FT /label= Phosphorylation
 FT /note= "potential casein-kinase 2 phosphorylation
 FT site"
 FT modified_site 939
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FT FT /note= "potential N-linked glycosylation site"
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Query Match 50.0%; Score 35; DB 1; Length 1784;
Best Local Similarity 26.7%; Pred. NO. 2.6e+02;
Matches 4; Conservative 7; Mismatches 4; Indels 0; Gaps 0;

QY 1 AYRLIKVIRIVLKY 15

DB 335 SYEIVLSITRLIKKY 349

RESULT 7

ID W76789 standard; Protein; 33 AA.

AC W76789;

DT 15-JAN-1999 (first entry)

DE D. immitis pDIAK1745 Ankyrin-like repeat protein fragment #12.

KW Ankyrin; helminth; parasite; vaccine; therapy; infection;

KV passive immunogen; cytotoxic agent.

OS Dirofilaria immitis.

PN US5827692-A.

PD 27-OCT-1998.

PF 24-APR-1997; 847429.

PR 24-APR-1997; US-847429.

PA (HESK-) HESKA CORP.

PI Biehm ES, Tang L;

DR WPI; 98-593992/50.

PT Nucleic acids encoding ankyrins from helminth parasites - useful for

recombinant production of the proteins for use as vaccines and

treatments against helminth infection

PS Example 2: Column 35-36; 84pp; English.

CC W76789-W76802 are ankyrin-like repeat peptide fragments isolated from the

helminth parasite Dirofilaria immitis pDIAK1075 protein. Such ankyrin

nucleic acids and recombinant products are useful for the recombinant

production of the ankyrin polypeptides. These proteins can then be used

as vaccines against parasitic helminth, e.g. D. immitis or B. malayi.

CC They can also be used for therapy after infection, and to raise

antibodies, also for use in therapeutics, as passive immunogens, or as

therapeutics against helminths on conjugation to cytotoxic agents. The

nucleic acids contained in viruses, may also be used as viral vaccines,

and the nucleic acids themselves or in vectors may be used as genetic

vaccines.

CC Sequence 33 AA;

Query Match 48.6%; Score 34; DB 1; Length 33;

Best Local Similarity 50.0%; Pred. NO. 6.1;

Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15

DB 15 IKIVELLKY 24

RESULT 8

ID W77378 standard; peptide; 37 AA.

AC W77378;

DT 14-DEC-1998 (first entry)

DE Lytic peptide with alterable function 3.
 KW Biologically active peptide; drug; toxin;
 KW lipid bilayer membrane; microorganism; parasite; virus.
 OS Synthetic.
 PN WO9841535-A2.
 PD 24-SEP-1998.
 PF 18-MAR-1998; G00799.
 PR 18-MAR-1997; GB-005519.
 PA (ANMA-) ANMAT TECHNOLOGY LTD.
 PI Ajoula HS, Clarke DJ;
 DR WPI; 98-521161/44.
 PT New modified peptide(s) - obtained by substitution with an amino
 PT acid which is modifiable by a reaction and replacing other amino
 PT acid which are not to be modified
 PS Claim 7; Page 22; 33pp; English.
 CC The peptides W77376-W77390 can be modified by the method of the
 CC invention by substituting at least one amino acid of the peptide to
 CC provide a peptide having at least one amino acid which is modifiable by a
 CC reaction and replacing other amino acids in the peptide with amino acids
 CC which are not modifiable by the reaction. The methods can be used for
 CC the modification of biologically active peptides such as hormones, drugs,
 CC toxins and peptides which act on lipid bilayer membranes. The modified
 CC peptides can be used e.g. in the body of an animal or plant or parts in
 CC order to affect the structure or integrity or permeability of a foreign
 CC body such as a microorganism, parasite or virus present in the body of
 CC the animal or plant or within the cells of the body of the animal or
 CC plant.
 SQ Sequence 37 AA;

Query Match 48.6%; Score 34; DB 1; Length 37;
 Best Local Similarity 41.7%; Pred. No. 6.9;
 Matches 5; Conservative 7; Mismatches 0; Indels 0; Gaps 0;

QY 3 RLLIKVIRVLK 14
 :||:|||||
 Db 4 KLLKLLKLLK 15

RESULT 9
 W76774
 ID W76774 standard; Protein; 303 AA.
 AC W76774;
 DT 15-JAN-1999 (first entry)
 DE D. immitis ankyrin pAnk303 protein.
 KW Ankyrin; helminth; parasite; vaccine; therapy; infection;
 KW passive immunogen; cytotoxic agent.
 OS Dirofilaria immitis.
 PN US5827692-A.
 PD 27-OCT-1998.
 PF 24-APR-1997; 847429.
 PR 24-APR-1997; US-847429.
 PA (HESK-) HESKA CORP.
 PI Blehm ES, Tang L;
 DR WPI; 98-593992/50.
 DR N-PSDB; V63012.
 PT Nucleic acids encoding ankyrins from helminth parasites - useful for
 PT recombinant production of the proteins for use as vaccines and
 PT treatments against helminth infection
 PS Claim 8; Column 79-82; 84pp; English.
 CC W76769-W76777 represent ankyrin proteins isolated from the helminth
 CC parasites Dirofilaria immitis and Brugia malayi. The nucleic acids and
 CC recombinant products are useful for the recombinant production of the
 CC ankyrin polypeptides. These proteins can then be used as vaccines against
 CC parasitic helminth, e.g. D. immitis or B. malayi. They can also be used
 CC for therapy after infection, and to raise antibodies, also for use in
 CC therapeutics, as passive immunogens, or as therapeutics against helminths
 CC on conjugation to cytotoxic agents. The nucleic acids contained in
 CC viruses, may also be used as viral vaccines, and the nucleic acids
 CC themselves or in vectors may be used as genetic vaccines.
 SQ Sequence 303 AA;

Query Match 48.6%; Score 34; DB 1; Length 303;
 Best Local Similarity 50.0%; Pred. No. 62;
 Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15
 |||:|||||
 Db 54 IKIVELLKY 63

RESULT 10
 W70606
 ID W70606 standard; Protein; 303 AA.
 AC W70606;
 DT 21-JAN-1999 (first entry)
 DE Ankyrin protein fragment pAnk303.
 KW Ankyrin protein; parasitic helminth; heartworm disease;
 KW elephantiasis; hydrocele.
 OS Dirofilaria immitis.
 PN US5824306-A.
 PD 20-OCT-1998.
 PF 26-FEB-1998; 031485.
 PR 24-APR-1997; US-847429.
 PR 26-FEB-1998; US-031485.
 PA (HESK-) HESKA CORP.
 PI Blehm ES, Tang L;
 DR WPI; 98-593373/50.
 DR N-PSDB; V63312.
 PT Dirofilaria and Brugia ankyrin proteins and antibodies - useful for
 PT protection of animals from disease caused by parasitic helminth
 PS Claim 5; Columns 79-82; 84pp; English.
 CC The present sequence represents part of a Dirofilaria immitis ankyrin
 CC protein. The ankyrin protein, or anti-ankyrin antibodies, may be used
 CC to protect an animal from disease caused by a parasitic helminth,
 CC especially where the disease is heartworm disease, elephantiasis or
 CC hydrocele.
 SQ Sequence 303 AA;

Query Match 48.6%; Score 34; DB 1; Length 303;
 Best Local Similarity 50.0%; Pred. No. 62;
 Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15
 |||:|||||
 Db 54 IKIVELLKY 63

RESULT 11
 W88282
 ID W88282 standard; Protein; 395 AA.
 AC W88282;
 DT 12-APR-1999 (first entry)
 DE Pseudomonas mendocina para-hydroxybenzoate hydroxylase.
 KW Para-hydroxybenzoate hydroxylase; para-hydroxybenzoic acid; PHBA;
 KW Poba-2.
 OS Pseudomonas mendocina.
 PN WO9856920-A1.
 PD 17-DEC-1998.
 PF 11-JUN-1998; U12072.
 PR 03-JUN-1998; US-049556.
 PR 13-JUN-1997; US-049556.
 PA (DUPO) DU PONT DE NEMOURS & CO E I.
 PI Chen KK, Grellak RL;
 DR WPI; 99-080332/05.
 DR N-PSDB; V84271.
 PT Production of para-hydroxybenzoate - using poba(-) Pseudomonas
 PT strains which have a toluene degradation pathway and do not produce
 PT para-hydroxybenzoate hydroxylase
 PS Claim 4; Page 39-41; 60pp; English.
 CC This is the amino acid sequence of a Pseudomonas mendocina
 CC para-hydroxybenzoate hydroxylase (PHB). P. mendocina possesses
 CC two highly homologous poba genes (see V84270-71), both of which
 CC encode PHB enzymes (see W88281-82) able to convert

CC para-hydroxybenzoic acid (PHBA) to protocatechuate. A claimed
 CC method for the production of PHBA comprises: (a) culturing a
 CC poba(-) Pseudomonas strain in a medium containing an aromatic
 CC organic substrate (e.g. toluene or p-cresol), at least one C-source
 CC (e.g. glucose or succinate) and an N-source, where the poba(-)
 CC Pseudomonas strain comprises poba genes encoding the toluene
 CC monooxygenase toluene degradation pathway and where the poba(-)
 CC Pseudomonas strain does not produce any detectable PHBH activity
 CC owing to poba gene disruption, so that PHBA accumulates at a rate
 CC of about 0.01-1 g PHBA/g cell hr; and (b) recovering the PHBA. The
 CC PHBA is used as a monomer for liquid crystal polymers. Esters of
 CC PHBA can also be used as backbone modifiers in condensation
 CC polymers, e.g. polyesters, and are also used to make paraben
 CC preservatives.
 CC Sequence 395 AA;

Query Match 48.6%; Score 34; DB 1; Length 395;
 Best Local Similarity 56.7%; Pred. No. 81;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 YRLIKVIR 10
 DB 310 YRLIVKVR 318

RESULT 12

ID P81184
 AC P81184 standard; Protein; 1594 AA.

DT 29-OCT-1992 (first entry)
 DE Sequence encoded by the 2nd reading frame of the peplomeric gene
 DE of FIVP strain 79-1146.
 KW Vaccine; peplomeric protein; E2 gene.
 OS Feline infectious peritonitis virus.
 PN EP-264979-A.

PD 27-APR-1988.

PF 01-SEP-1987; 201657.

PR 05-SEP-1986; NL-002244.

PA (DUIN) DUPHAR INT RES BV.

PI De Groot RJ, Spaan WJM, Van Der Zeijst BAM;

DR WPI: 88-114147/17.

DR N-PSDB; N81533.

PT Gene for feline infectious peritonitis virus - and gene prod.

PT useful as antigenic protein for vaccine

PS Disclosure; Fig 1; 13pp; English.

CC cDNA was prep'd from FIVP strain 79-1146. N81533 gives the sequence

CC of the peplomeric gene in three reading frames. The top reading

CC frame is an open reading frame of 4356 nucleotides and has a coding

CC capacity for a precursor polypeptide having a mol. wt. of 160,470

CC (1452 AAs). The beginning and the end of the E2 gene are indicated

CC in the FT of N81533. The first 18 N-terminal AAs have a strong

CC hydrophobic character and presumably comprise a cleavable signal

CC peptide. The extreme carboxy-terminal part comprises a region of 20

CC hydrophobic AAs, which presumably serves as a transmembrane anchor.

CC The FIVP peplomeric protein has 35 potential glycosylation sites,

CC of which 22 are in the N-terminal part (pos. 1-790) which corresponds

CC to the S-part of the IBV E2 (see P81183). N.B. IBV = infectious

CC bronchitis virus. "X" in the AA sequence denotes the translation

CC of a stop codon.

CC Sequence 1594 AA;

Query Match 48.6%; Score 34; DB 1; Length 1594;

Best Local Similarity 57.1%; Pred. No. 3.5e+02;

Matches 8; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 2 YRLIKVIRVILKY 15

DB 787 YRLIIAVCTICLCY 800

RESULT 13

W76776

ID W76776 standard; Protein; 1745 AA.

AC W76776;

DT 15-JAN-1999 (first entry)

DE D. immitis ankyrin p1Ank1745 protein.

KW Ankyrin; helminth; parasite; vaccine; therapy; infection;

OS passive immunogen; cytotoxic agent.

OS Dirofilaria immitis.

PN US5827692-A.

PD 27-OCT-1998.

PF 24-APR-1997; 847429.

PR 24-APR-1997; US-847429.

PA (HESK-) HESKA CORP.

PI Blehm ES, Tang L;

DR WPI: 98-593992/50.

DR N-PSDB; V63020.

PT Nucleic acids encoding ankyrins from helminth parasites - useful for

PT recombinant production of the proteins for use as vaccines and

PT treatments against helminth infection

PS Claim 8; Column 109-118; 84pp; English.

CC W76769-W76777 represent ankyrin proteins isolated from the helminth

CC parasites Dirofilaria immitis and Brugia malayi. The nucleic acids and

CC recombinant products are useful for the recombinant production of the

CC ankyrin polypeptides. These proteins can then be used as vaccines against

CC parasitic helminth, e.g. D. immitis or B. malayi. They can also be used

CC for therapy after infection, and to raise antibodies, also for use in

CC therapeutics, as passive immunogens, or as therapeutics against helminths

CC on conjugation to cytotoxic agents. The nucleic acids contained in

CC viruses, may also be used as viral vaccines, and the nucleic acids

CC themselves or in vectors may be used as genetic vaccines.

CC Sequence 1745 AA;

Query Match 48.6%; Score 34; DB 1; Length 1745;

Best Local Similarity 50.0%; Pred. No. 3.8e+02;

Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15

DB 393 IKIVELLKY 402

RESULT 14

ID W70608

AC W70608 standard; Protein; 1745 AA.

DT 21-JAN-1999 (first entry)

DE Full length ankyrin protein.

KW Ankyrin protein; parasitic helminth; heartworm disease;

OS elephantiastis; hydrocele.

OS Dirofilaria immitis.

PN US5824306-A.

PD 20-OCT-1998.

PF 26-FEB-1998; 031485.

PR 24-APR-1997; US-847429.

PR 26-FEB-1998; US-031485.

PA (HESK-) HESKA CORP.

PI Blehm ES, Tang L;

DR WPI: 98-593373/50.

DR N-PSDB; V63314.

PT Dirofilaria and Brugia ankyrin proteins and antibodies - useful for

PT protection of animals from disease caused by parasitic helminth

PS Claim 5; Columns 107-118; 84pp; English.

CC The present sequence represents a full length dirofilaria immitis ankyrin

CC protein. The ankyrin protein, or anti-ankyrin antibodies, may be used

CC to protect an animal from disease caused by a parasitic helminth, or

CC especially where the disease is heartworm disease, elephantiastis or

CC hydrocele.

CC Sequence 1745 AA;

Query Match 48.6%; Score 34; DB 1; Length 1745;

Best Local Similarity 50.0%; Pred. No. 3.8e+02;

Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15

|||||

Db 393 IKIVELLKY 402

RESULT 15

R92907
ID R92907 standard; peptide; 20 AA.

AC R92907:

DT 16-MAY-1996' (first entry)

DE HLA-B2702 CTL modulating peptide (B2702.84-75/75-84).

KW Cytotoxic T lymphocyte; CTL; major histocompatibility complex; MHC;

KW immunosuppressant; graft versus host disorder; transplantation; therapy;

KW class I MHC; HLA-B2702.

OS Synthetic.

PN W09526979-A1.

PD 12-OCT-1995.

PF 05-APR-1995; U04349.

PR 05-APR-1994; US-222851.

PA (STRD) UNIV LELAND STANFORD JUNIOR.

PI Clayberger C, Krensky AM, Parham P;

DR WPI; 95-358582/46.

PT Extension of acceptance period of transplants from MHC unmatched

PT donor hosts; - using Class I B*5-84 MHC antigen of the recipient

PT host

PS Example 15; Page 36; 80pp; English.

CC R83061-R83085, R83090-R83096 and R92907-R92914 represent fragments of

CC class I major histocompatibility complex (MHC) antigens. This sequence

CC is an inverted dimer of residues 75-84 of the alpha-1 domain of the class

CC I MHC HLA-B2702. These sequences can be used to extend the period of

CC acceptance by a recipient of a transplant from an MHC unmatched donor.

CC The peptides are administered to a patient in conjunction with a

CC subtherapeutic amount of an immunosuppressant. This is administered to

CC the patient for a limited period of time (compared to the lifetime

CC administration for current treatments). The peptides particularly

CC modulate (or inhibit) the activity of the cytotoxic T lymphocytes (CTLs)

CC of the patient.

SQ Sequence 20 AA;

Query Match

Best Local Similarity 47.1%; Score 33; DB 1; Length 20;

Matches 8; Conservative 4; Mismatches 2; Indels 6; Gaps 1;

QY 2 YRLIKV-----IRIVLKY 15

|||||

Db 1 YRLAIRNERERLRIALRY 20

Search completed: February 8, 2000, 04:05:40

Job time: 9337 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 00:02:05 ; Search time 75.85 Seconds
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2.541 Million cell updates/sec

Title: US-08-653-294-21

Perfect score: 70
Sequence: 1 AYRLIKVIRVLKY 15

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 130275 seqs, 12848600 residues

Total number of hits satisfying chosen parameters: 130275

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query	Score	Match	Length	ID	Description
1	36	51.4	222	2	US-08-756-771-5	Sequence 5, Appli
2	36	51.4	222	2	US-09-096-571-5	Sequence 5, Appli
3	36	51.4	1226	2	US-08-540-804-12	Sequence 12, Appl
4	36	51.4	1226	2	US-08-218-265-12	Sequence 12, Appl
5	35	50.0	302	2	US-09-031-485-38	Sequence 38, Appl
6	35	50.0	302	2	US-08-847-429A-38	Sequence 38, Appl
7	34	48.6	33	2	US-09-031-485-72	Sequence 72, Appl
8	34	48.6	33	2	US-08-847-429A-72	Sequence 72, Appl
9	34	48.6	303	2	US-09-031-485-23	Sequence 23, Appl
10	34	48.6	303	2	US-08-847-429A-23	Sequence 23, Appl
11	34	48.6	1745	2	US-09-031-485-33	Sequence 33, Appl
12	34	48.6	1745	2	US-08-847-429A-33	Sequence 33, Appl
13	33	47.1	20	1	US-08-222-851-33	Sequence 33, Appl
14	33	47.1	20	3	PCT-US94-12985-1	Sequence 1, Appli
15	33	47.1	20	3	PCT-US94-12985-4	Sequence 4, Appli
16	33	47.1	2860	2	US-08-826-267-2	Sequence 2, Appli
17	32	45.7	12	3	PCT-US94-12985-5	Sequence 5, Appli
18	32	45.7	227	1	US-08-597-236-6	Sequence 6, Appli
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20	32	45.7	384	2	US-08-637-759B-375	Sequence 375, App
21	32	45.7	575	2	US-08-766-858A-5	Sequence 5, Appli
22	31	44.3	101	1	US-07-820-154A-7	Sequence 7, Appli
23	31	44.3	101	1	US-08-220-401-5	Sequence 5, Appli
24	31	44.3	101	2	US-08-437-362-5	Sequence 5, Appli
25	31	44.3	101	2	US-08-097-554A-7	Sequence 7, Appli
26	31	44.3	101	3	PCT-US93-00324-7	Sequence 7, Appli
27	31	44.3	437	2	US-08-538-816A-2	Sequence 2, Appli
28	31	44.3	437	2	US-09-076-651-2	Sequence 2, Appli
29	30	42.9	27	1	US-08-343-427B-1	Sequence 1, Appli
30	30	42.9	27	1	US-08-343-427B-2	Sequence 2, Appli

Sequence 3, Appli
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37 30 42.9 32 2 US-08-652-450A-4
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42 30 42.9 35 4 5223481-5
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ALIGNMENTS

RESULT 1
US-08-756-771-5
; Sequence 5, Application US/08756771
; Patent No. 5817497
; GENERAL INFORMATION:
; APPLICANT: Goli, Surya K.
; APPLICANT: Hillman, Jennifer L.
; TITLE OF INVENTION: A NOVEL GLUTATHIONE S-TRANSFERASE
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: US
; ZIP: 94304

COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: DOS
SOFTWARE: FastSeq Version 2.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/756,771
FILING DATE: Herewith
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER:
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Billings, Lucy J.
REGISTRATION NUMBER: 36,749
REFERENCE/DOCKET NUMBER: PF-0162 US
TELECOMMUNICATION INFORMATION:
TELEPHONE: 415-855-0555
TELEFAX: 415-845-4166
TELEX:

INFORMATION FOR SEQ ID NO: 5:
SEQUENCE CHARACTERISTICS:
LENGTH: 222 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
IMMEDIATE SOURCE:
LIBRARY: GenBank
CLONE: 193710

US-08-756-771-5
Query Match 51.4%; Score 36; DB 2; Length 222;
Best Local Similarity 60.0%; Pred No. 16;
Matches 6; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 6 IKVIRVLKY 15

Db 213 VEVVRLVKF 222
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RESULT 2

US-09-096-571-5
; Sequence 5, Application US/09096571
; Patent No. 5976528
; GENERAL INFORMATION:
; APPLICANT: Goli, Surya K.
; APPLICANT: Hillman, Jennifer L.
; TITLE OF INVENTION: A NOVEL GLUTATHIONE S-TRANSFERASE
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: INCYTE PHARMACEUTICALS, INC.
; STREET: 3174 Porter Drive
; CITY: Palo Alto
; STATE: CA
; COUNTRY: US
; ZIP: 94304
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FASTSEQ Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/096.571
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/756,771
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Billings, Lucy J.
; REGISTRATION NUMBER: 36,749
; REFERENCE/DOCKET NUMBER: PF-0162 US
; TELEPHONE: 415-855-0555
; TELEFAX: 415-845-4166
; TELEX:
; INFORMATION FOR SEQ ID NO: 5:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 222 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; LIBRARY: GenBank
; CLONE: 193710
; US-09-096-571-5

Query Match 51.4%; Score 36; DB 2; Length 222;
Best Local Similarity 60.0%; Pred. No. 16;
Matches 6; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

Qy 6 IKVIRVLKY 15
:::|||||

Db 213 VEVVRLVKF 222

RESULT 3

US-08-540-804-12
; Sequence 12, Application US/08540804
; Patent No. 5919666
; GENERAL INFORMATION:
; APPLICANT: Young, Richard A.
; APPLICANT: Koleske, Anthony J.
; APPLICANT: Thompson, Craig M.
; APPLICANT: Chao, David M.
; TITLE OF INVENTION: No. 5919666el Factors Which Modify Gene
; NUMBER OF SEQUENCES: 39

; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: Massachusetts
; COUNTRY: US
; ZIP: 02173
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/540,804
; FILING DATE: 11-OCT-1995
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/521,872
; FILING DATE: 21-AUG-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/218,265
; FILING DATE: 25-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Granahan, Patricia
; REGISTRATION NUMBER: 32,227
; REFERENCE/DOCKET NUMBER: WH194-03A2
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 617-861-6240
; TELEFAX: 617-861-9540
; INFORMATION FOR SEQ ID NO: 12:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1226 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-540-804-12

Query Match 51.4%; Score 36; DB 2; Length 1226;
Best Local Similarity 77.8%; Pred. No. 90;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 YRLIKVIR 10
| |||||

Db 999 YHLLIKIIR 1007

RESULT 4

US-08-218-265-12
; Sequence 12, Application US/08218265
; Patent No. 5922585
; GENERAL INFORMATION:
; APPLICANT: Young, Richard A.
; APPLICANT: Koleske, Anthony J.
; APPLICANT: Thompson, Craig M.
; TITLE OF INVENTION: No. 5922585el Factors Which Modify Gene
; NUMBER OF SEQUENCES: 35
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: MA
; COUNTRY: US
; ZIP: 02173
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/218,265
; FILING DATE: 25-MAR-1994

CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Granahan, Patricia
REGISTRATION NUMBER: 32,227
REFERENCE/DOCKET NUMBER: WHI94-03
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-861-6240
TELEFAX: 617-861-9540
INFORMATION FOR SEQ ID NO: 12:
SEQUENCE CHARACTERISTICS:
LENGTH: 1226 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-218-265-12

Query Match 51.4%; Score 36; DB 2; Length 1226;
Best Local Similarity 77.8%; Pred. No. 90;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 YRLIKVIR 10
| |||||
Db 999 YHLLIKIIR 1007

RESULT 5
US-09-031-485-38
; Sequence 38, Application US/09031485
; Patent No. 5824306
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Carol Talkington Verser, Ph.D.
; ADDRESSEE: Heska Corporation
; STREET: 1825 Sharp Point Drive
; CITY: Fort Collins
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80525
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect for Windows, Version 7.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/031,485
; FILING DATE:
; CLASSIFICATION: 530
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/847,429
; FILING DATE: 24-APR-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Verser, Carol Talkington
; REGISTRATION NUMBER: 37,459
; REFERENCE/DOCKET NUMBER: HW-5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 970/493-7272
; TELEFAX: 970/484-9505
; INFORMATION FOR SEQ ID NO: 38:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 302 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-031-485-38

Query Match 50.0%; Score 35; DB 2; Length 302;
Best Local Similarity 60.0%; Pred. No. 33;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
QY 6 IKVIRIVLKY 15
| ||| : |||
Db 53 IKVVELLKY 62
| ||| : |||
RESULT 6
US-08-847-429A-38
; Sequence 38, Application US/08847429A
; Patent No. 5827692
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Carol Talkington Verser, Ph.D.
; ADDRESSEE: Heska Corporation
; STREET: 1825 Sharp Point Drive
; CITY: Fort Collins
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80525
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect for Windows, Version 7.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/847,429A
; FILING DATE: 24-APR-1997
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Verser, Carol Talkington
; REGISTRATION NUMBER: 37,459
; REFERENCE/DOCKET NUMBER: HW-5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 970/493-7272
; TELEFAX: 970/484-9505
; INFORMATION FOR SEQ ID NO: 38:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 302 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-847-429A-38

Query Match 50.0%; Score 35; DB 2; Length 302;
Best Local Similarity 60.0%; Pred. No. 33;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;
QY 6 IKVIRIVLKY 15
| ||| : |||
Db 53 IKVVELLKY 62
| ||| : |||

RESULT 7
US-09-031-485-72
; Sequence 72, Application US/09031485
; Patent No. 5824306
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85

;; CORRESPONDENCE ADDRESS:
;; ADDRESSEE: Carol Talkington Verser, Ph.D.
;; ADDRESSEE: Heskia Corporation
;; STREET: 1825 Sharp Point Drive
;; CITY: Fort Collins
;; STATE: Colorado
;; COUNTRY: USA
;; ZIP: 80525
;; COMPUTER READABLE FORM:
;; MEDIUM TYPE: Floppy disk
;; COMPUTER: IBM PC compatible
;; OPERATING SYSTEM: Windows 95
;; SOFTWARE: WordPerfect for Windows, Version 7.0
;; CURRENT APPLICATION DATA:
;; APPLICATION NUMBER: US/09/031,485
;; FILING DATE:
;; CLASSIFICATION: 530
;; PRIOR APPLICATION DATA:
;; APPLICATION NUMBER: US/08/847,429
;; FILING DATE: 24-APR-1997
;; ATTORNEY/AGENT INFORMATION:
;; NAME: Verser, Carol Talkington
;; REGISTRATION NUMBER: 37,459
;; REFERENCE/DOCKET NUMBER: HW-5
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 970/493-7272
;; TELEFAX: 970/484-9505
;; INFORMATION FOR SEQ ID NO: 72:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 33 amino acids
;; TYPE: amino acid
;; TOPOLOGY: linear
;; MOLECULE TYPE: protein
;; US-09-031-485-72

Query Match 48.6%; Score 34; DB 2; Length 33;
Best Local Similarity 50.0%; Pred. No. 5.2;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15
Db 15 IKIVELLKY 24

RESULT 8
US-08-847-429A-72
; Sequence 72, Application US/08847429A
; Patent No. 5827692
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Carol Talkington Verser, Ph.D.
; ADDRESSEE: Heskia Corporation
; STREET: 1825 Sharp Point Drive
; CITY: Fort Collins
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80525
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect for Windows, Version 7.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/847,429A
; FILING DATE: 24-APR-1997
; CLASSIFICATION: 435

;; ATTORNEY/AGENT INFORMATION:
;; NAME: Verser, Carol Talkington
;; REGISTRATION NUMBER: 37,459
;; REFERENCE/DOCKET NUMBER: HW-5
;; TELECOMMUNICATION INFORMATION:
;; TELEPHONE: 970/493-7272
;; TELEFAX: 970/484-9505
;; INFORMATION FOR SEQ ID NO: 72:
;; SEQUENCE CHARACTERISTICS:
;; LENGTH: 33 amino acids
;; TYPE: amino acid
;; TOPOLOGY: linear
;; MOLECULE TYPE: protein
;; US-08-847-429A-72

Query Match 48.6%; Score 34; DB 2; Length 33;
Best Local Similarity 50.0%; Pred. No. 5.2;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

QY 6 IKVIRIVLKY 15
Db 15 IKIVELLKY 24

RESULT 9
US-09-031-485-23
; Sequence 23, Application US/09031485
; Patent No. 5824306
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Carol Talkington Verser, Ph.D.
; ADDRESSEE: Heskia Corporation
; STREET: 1825 Sharp Point Drive
; CITY: Fort Collins
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80525
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect for Windows, Version 7.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/031,485
; FILING DATE:
; CLASSIFICATION: 530
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/847,429
; FILING DATE: 24-APR-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Verser, Carol Talkington
; REGISTRATION NUMBER: 37,459
; REFERENCE/DOCKET NUMBER: HW-5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 970/493-7272
; TELEFAX: 970/484-9505
; INFORMATION FOR SEQ ID NO: 23:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 303 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-031-485-23

Query Match 48.6%; Score 34; DB 2; Length 303;

Best Local Similarity 50.08; Pred. No. 49; Indels 1; Gaps 0;
Matches 5; Conservative 4; Mismatches 0;

Qy 6 IKVIRVLKY 15
|::: :|||
Db 54 IKIVELLKY 63

RESULT 10
US-08-847-429A-23
; Sequence 23, Application US/08847429A
; Patent No. 5827692
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Carol Talkington Verser, Ph.D.
; ADDRESSEE: Heska Corporation
; STREET: 1825 Sharp Point Drive
; CITY: Fort Collins
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80525

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect for Windows, Version 7.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/847,429A
FILING DATE: 24-APR-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Verser, Carol Talkington
REGISTRATION NUMBER: 37,459
REFERENCE/DOCKET NUMBER: HW-5
TELECOMMUNICATION INFORMATION:
TELEPHONE: 970/493-7272
TELEFAX: 970/484-9505
INFORMATION FOR SEQ ID NO: 23:
LENGTH: 303 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-847-429A-23

Query Match 48.68; Score 34; DB 2; Length 303;
Best Local Similarity 50.08; Pred. No. 49; Indels 1; Gaps 0;
Matches 5; Conservative 4; Mismatches 0;

Qy 6 IKVIRVLKY 15
|::: :|||
Db 54 IKIVELLKY 63

RESULT 11
US-09-031-485-33
; Sequence 33, Application US/09031485
; Patent No. 5824306
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:

ADDRESSEE: Carol Talkington Verser, Ph.D.
ADDRESSEE: Heska Corporation
STREET: 1825 Sharp Point Drive
CITY: Fort Collins
STATE: Colorado
COUNTRY: USA
ZIP: 80525
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect for Windows, Version 7.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/031,485

FILING DATE:
CLASSIFICATION: 530
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/847,429
FILING DATE: 24-APR-1997
ATTORNEY/AGENT INFORMATION:
NAME: Verser, Carol Talkington
REGISTRATION NUMBER: 37,459
REFERENCE/DOCKET NUMBER: HW-5
TELECOMMUNICATION INFORMATION:
TELEPHONE: 970/493-7272
TELEFAX: 970/484-9505
INFORMATION FOR SEQ ID NO: 33:
LENGTH: 1745 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-031-485-33

Query Match 48.68; Score 34; DB 2; Length 1745;
Best Local Similarity 50.08; Pred. No. 2.9e+02;
Matches 5; Conservative 4; Mismatches 1; Indels 0; Gaps 0;

Qy 6 IKVIRVLKY 15
|::: :|||
Db 393 IKIVELLKY 402

RESULT 12
US-08-847-429A-33
; Sequence 33, Application US/08847429A
; Patent No. 5827692
; GENERAL INFORMATION:
; APPLICANT: Tang, Liang
; APPLICANT: Blehm, E. Scot
; TITLE OF INVENTION: DIROFILARIA AND BRUGIA ANKYRIN
; TITLE OF INVENTION: PROTEINS, NUCLEIC ACID MOLECULES, AND
; TITLE OF INVENTION: USES THEREOF
; NUMBER OF SEQUENCES: 85
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Carol Talkington Verser, Ph.D.
; ADDRESSEE: Heska Corporation
; STREET: 1825 Sharp Point Drive
; CITY: Fort Collins
; STATE: Colorado
; COUNTRY: USA
; ZIP: 80525

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect for Windows, Version 7.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/847,429A
FILING DATE: 24-APR-1997
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:

Qv 2 YRLIKV-----IRIVLY 15

PC/US94-12985-4
 ; Sequence 4, Application PC/TUS9412985
 ; GENERAL INFORMATION:
 ; APPLICANT: The Board of Trustees for the Leland Stanford Junior
 ; APPLICANT: University
 ; TITLE OF INVENTION: SURFACE MEMBRANE PROTEINS AND THEIR
 ; TITLE OF INVENTION: EFFECT ON IMMUNE RESPONSE
 ; NUMBER OF SEQUENCES: 20
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSSEE: FLEHR, HOHBACH, TEST, ALBRITTON & HERBERT
 ; STREET: 4 Embarcadero Center, Suite 3400
 ; CITY: San Francisco
 ; STATE: California

```

: COUNTRY: USA
: ZIP: 94111-4187
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patent In Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: PCT/US94/12985
: FILING DATE: 10-NOV-1994
: CLASSIFICATION:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: 08/150,493
: FILING DATE: 10-NOV-1993
: ATTORNEY/AGENT INFORMATION:
: NAME: Rowland, Bertram I
: REGISTRATION NUMBER: 20,015
: REFERENCE/DOCKET NUMBER: FP-58976-PC/BIR
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (415) 781-1989
: TELEFAX: (415) 398-3249
: TELEX: 910 277299
: INFORMATION FOR SEQ ID NO: 4:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 20 amino acids
: TYPE: amino acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: peptide
: PCT-US94-12985-4

```

```

Query Match      47.1%; Score 33; DB 3; Length 20;
Best Local Similarity 40.0%; Pred. No. 4.7;
Matches 8; Conservative 4; Mismatches 2; Indels 6; Gaps 1;

QY  2 YRLIKV-----IRVLKY 15
    ||| |::: ||| |::|
Db   1 YRLAIRLNERNLRY 20

```

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Search completed: February 8, 2000, 01:01:46
Job time: 3581 sec

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 7, 2000, 11:54:34 ; Search time 111.22 Seconds
(without alignments)
6.362 Million cell updates/sec

Title: US-08-653-294-21

Perfect score: 70

Sequence: 1 AYRLIKVIRVLKY 15

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 142080 seqs, 47169319 residues

Total number of hits satisfying chosen parameters: 142080

Minimum DB seq length: 0

Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%

Listing first 45 summaries

Database : PIR.62:*

1: pir1:*

2: pir2:*

3: pir3:*

4: pir4:*

pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	39.5	56.4	746	2 S67203	probable membrane
2	38	54.3	404	2 F64238	hypothetical prote
3	38	54.3	709	1 OXCKPM	acyl-CoA oxidase (
4	38	54.3	771	2 T16254	hypothetical prote
5	37	52.9	98	2 C32227	hypothetical prote
6	37	52.9	242	2 G75029	hypothetical prote
7	37	52.9	379	2 G75170	hypothetical prote
8	36.5	52.1	724	2 S57604	probable membrane
9	36	51.4	131	2 G70326	hypothetical prote
10	36	51.4	186	2 C71601	probable integral
11	36	51.4	187	2 E71464	probable rRNA meth
12	36	51.4	222	2 S27334	glutathione trans
13	36	51.4	282	2 B69260	hypothetical prote
14	36	51.4	432	2 S65238	probable membrane
15	36	51.4	705	2 S38066	probable finger pr
16	36	51.4	1427	2 S74293	SRB8 protein - vea
17	36	51.4	2048	1 ZLN2SE	genome polyprotein
18	36	51.4	2228	1 ZLN2SE	genome polyprotein
19	35	50.0	341	2 A72674	hypothetical prote
20	35	50.0	485	2 T09374	H+-transporting Ar
21	35	50.0	486	2 B70775	probable atpD prot
22	35	50.0	502	1 OXCKAX	acyl-CoA oxidase (
23	35	50.0	709	1 OXCKX4	acyl-CoA oxidase (
24	35	50.0	709	1 OXCKX	acyl-CoA oxidase (
25	35	50.0	791	2 T02583	hypothetical prote
26	35	50.0	1784	2 A49420	tuberos sclerosi
27	35	50.0	1786	2 A57282	ankyrin-related pr
28	35	50.0	1809	2 T15345	ankyrin-related un
29	35	50.0	1809	2 S57329	tuberos sclerosi
30	35	50.0	1815	2 T15346	elegans ankyrin-re

31	35	50.0	1867	2 T15344	ankyrin-related un
32	35	50.0	2039	2 T15347	ankyrin-related un
33	35	50.0	2376	2 S48405	probable membrane
34	35	50.0	3973	2 B71612	hypothetical prote
35	34	48.6	180	2 G72616	hypothetical prote
36	34	48.6	185	2 D72118	hypothetical prote
37	34	48.6	277	2 JC2365	replication protei
38	34	48.6	326	2 S59101	NADH dehydrogenase
39	34	48.6	359	2 H71343	hypothetical prote
40	34	48.6	369	2 T15213	hypothetical prote
41	34	48.6	376	2 H71689	probable UDP-n-ace
42	34	48.6	395	2 S74051	hypothetical prote
43	34	48.6	448	2 S56260	probable membrane
44	34	48.6	651	2 G71697	probable soluble 1
45	33	47.1	111	2 A72079	hypothetical prote

ALIGNMENTS

RESULT 1
S67203
probable membrane protein YOR299w - yeast (Saccharomyces cerevisiae)
N:Alternate names: hypothetical protein O5635
C:Species: Saccharomyces cerevisiae
C>Date: 12-Jul-1996 #sequence_revision 12-Jul-1996 #text_change 06-Feb-1998
C:Accession: S67203
R:Cziepluch, C.; Jauniaux, J.C.; Kordes, E.; Poirey, R.; Pujol, A.; Tobiasch, E.
submitted to the Protein Sequence Database, July 1996
A:Reference number: S67194
A:Accession: S67203
A:Molecule type: DNA
A:Residues: 1-746 <GZI>
A:Cross-references: EMBL:275207; NID:g1420661; PID:e252136; PID:g1420662; MIPS:YOR299
A:Experimental source: strain S288C
C:Genetics:
A:Gene: SGD:BUD7
A:Cross-references: SGD:S0005825; MIPS:YOR299w
A:Map position: 15R
C:Keywords: transmembrane protein
F:328-344/Domain: transmembrane #status predicted <TM>

Query Match 56.4%; Score 39.5; DB 2; Length 746;
Best Local Similarity 45.0%; Pred. No. 17;
Matches 9; Conservative 5; Mismatches 1; Indels 5; Gaps 1;

QY 1 AYRLIKVIRI-----VLKY 15
||||| :|||
DB 502 AYRLTEIVQITGWENLLKY 521

RESULT 2
F64238
hypothetical protein MG349 - Mycoplasma genitalium (SGC3)
C:Species: Mycoplasma genitalium
C>Date: 10-Nov-1995 #sequence_revision 10-Nov-1995 #text_change 10-Oct-1997
C:Accession: F64238
R:Fraser, C.M.; Gocayne, J.D.; White, O.; Adams, M.D.; Clayton, R.A.; Fleischmann, R.
M.; Fuhrmann, J.; Nguyen, D.; Utterback, T.R.; Saudek, D.M.; Phillips, C.A.; Merrick,
C.A.; Venter, J.C.
Science 270, 397-403, 1995
A:Title: The minimal gene complement of Mycoplasma genitalium.
A:Reference number: A64200; MUID:96026346
A:Accession: F64238
A>Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-404 <TIGR>
A:Cross-references: GB:U39719; GB:L43967; NID:g1046055; PID:g1046056; TIGR:MG349
A:Experimental source: strain G-37
C:Genetics:
A:Genetic code: SGC3


```

Query Match          51.4%; Score 36; DB 2; Length 187;
Best Local Similarity 42.9%; Pred. No. 21;
Matches 6; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

OY 2 YRLIKRIVRLKY 15
      | : | : |||
      | : | : |||
Db 130 YNYLVNAITVWLKY 143

RESULT 12
S27234
glutathione transferase (EC 2.5.1.18) 5.7 - mouse
C:Species: Mus musculus (house mouse)
C:Date: 22-Nov-1993 #sequence_revision 10-Nov-1995 #text_change 18-Jun-1999
C:Accession: S27234; S17542; S17681
R:Zimniak, P.; Eckles, M.A.; Saxena, M.; Awasthi, Y.C.
FEBS Lett. 313, 173-176, 1992
A:A:Title: A subgroup of class alpha glutathione S-transferases. Cloning of cDNA for mo
A:Reference number: S27234; MUID:93050245
A:Accession: S27234
A:Molecule type: mRNA
A:Residues: 1-222 <ZIM>
A:Cross-references: GB:L06047; NID:g193709; PIDN:AAA37754.1; PID:g193710
R:R.Nedh, R.D.; Saxena, M.; Singhal, S.S.; Ahmad, H.; Awasthi, Y.C.
Biochem. J. 278, 793-799, 1991
A:A:Title: Characterization of a novel glutathione S-transferase isoenzyme from mouse l
A:Reference number: S17542; MUID:91378941
A:Accession: S17542
A:Molecule type: protein

```

Biocentre. J. 2/8, 793-799, 1991
 A:Title: Characterization of a novel glutathione S-transferase isoenzyme from mouse liver
 A:Reference number: SI7542; MUID:91378941
 A:Accession: SI7542
 A:Molecule type: protein

A:Residues: 106-114,'P',116-120 <MED>
A:Experimental source: lung
A:Accession: S17681

A:Molecule type: protein
A:Residues: 107-113,'G',168-178,'GE',181-184,'X',186-186 <MEW>
A:Experimental source: liver
C:Superfamily: glutathione transferase
C:Keywords: transferase

Query Match 51.4%; Score 36; DB 2; Length 222;
Best Local Similarity 60.0%; Pred. No. 25;
Matches 6; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 6 IKVIRVLKY 15

DB 213 VEVIRVLKF 222

RESULT 13

B9260
hypothetical protein AF0082 - Archaeoglobus fulgidus
C:Species: Archaeoglobus fulgidus
C:Date: 05-Dec-1997 #sequence_revision 05-Dec-1997 #text_change 05-Jun-1998
C:Accession: B9260
R:Klenk, H.P.; Clayton, R.A.; Tomb, J.F.; White, O.; Nelson, K.E.; Ketchum, K.A.; Dodson
.. Fleischmann, R.D.; Quackenbush, J.; Lee, N.H.; Sutton, G.G.; Gill, S.; Kirkness, E.F.;
Glodek, A.; Zhou, L.; Overbeek, R.; Gocayne, J.D.; Weidman, J.F.; McDonald, L.
Nature 390, 364-370, 1997
A:Authors: Utterback, T.; Cotton, M.D.; Spriggs, T.; Artach, P.; Kaine, B.P.; Sykes, S.
Smith, H.O.; Woese, C.R.; Venter, J.C.
A:Title: The complete genome sequence of the hyperthermophilic, sulfate-reducing archaeo
A:Reference number: A9250; MUID:98049343
A:Accession: B9260
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-282 <KLE>
A:Cross-references: GB:AE001100; GB:AE000782; NID:g2689423; PID:g2650567; TIGR:AF0082

Query Match 51.4%; Score 36; DB 2; Length 282;
Best Local Similarity 53.8%; Pred. No. 31;
Matches 7; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 2 YRLIKVIRVLK 14

DB 269 YRHLRLIRMTLK 281

RESULT 14

S65238
probable membrane protein YPL219w - yeast (Saccharomyces cerevisiae)
N:Alternate names: hypothetical protein P1745
C:Species: Saccharomyces cerevisiae
C:Date: 10-Dec-1994 #sequence_revision 31-May-1996 #text_change 06-Feb-1998
C:Accession: S65238
R:Rieger, M.; Mueller-Auer, S.; Schaefer, M.
submitted to the Protein Sequence Database, May 1996
A:Reference number: S65202
A:Accession: S65238
A:Molecule type: DNA
A:Residues: 1-492 <RIE>
A:Cross-references: EMBL:273575; NID:g1370453; PID:g246935; PID:g1370454; MIPS:YPL219w
A:Experimental source: strain S288C (AB972)
C:Genetics:
A:Gene: SGD:PC18
A:Cross-references: SGD:S0006140; MIPS:YPL219w
A:Map position: 16L
C:Keywords: transmembrane protein
F:380-396/Domain: transmembrane #status predicted <TMM>

Query Match 51.4%; Score 36; DB 2; Length 492;

Best Local Similarity 50.0%; Pred. No. 51;
Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 1 AYRLIKVIRVLK 14

DB 416 AHRIITIRIATK 429

RESULT 15

S38066
probable finger protein YKL222c - yeast (Saccharomyces cerevisiae)
N:Alternate names: hypothetical protein F705
C:Species: Saccharomyces cerevisiae
C:Date: 03-May-1994 #sequence_revision 03-May-1994 #text_change 24-Sep-1999
C:Accession: S38066; S4510; S43548
R:Alexandraki, D.; Horaitis, O.; Tzermia, M.
submitted to the Protein Sequence Database, March 1994
A:Reference number: S38065
A:Accession: S38066
A:Molecule type: DNA
A:Residues: 1-705 <ALE>
A:Cross-references: EMBL:Z28222; NID:9486398; PIDN:CAA82067.1; PID:9486399; MIPS:YKL2
A:Experimental source: strain S288C
R:Alexandraki, D.; Tzermia, M.
Yeast 10(Suppl.A), S81-S91, 1994
A:Title: Sequencing of a 13.2 kb segment next to the left telomere of yeast chromosom
V.
A:Reference number: S44508; MUID:94378726
A:Accession: S44510
A:Status: translation not shown
A:Molecule type: DNA
A:Residues: 1-705 <AL2>
A:Cross-references: EMBL:X75950; NID:g473124; PIDN:CAA53551.1; PID:g473127
A:Experimental source: strain S288C
C:Genetics:
A:Map position: 11L
C:Superfamily: unassigned GAL4-type zinc cluster proteins; GAL4 zinc binuclear cluste
C:Keywords: DNA binding; nucleus; zinc finger
F:19-57/Domain: GAL4 zinc binuclear cluster homology <GAL4>
F:24-52/Region: zinc finger

Query Match 51.4%; Score 36; DB 2; Length 705;
Best Local Similarity 46.7%; Pred. No. 71;
Matches 7; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

QY 1 AYRLIKVIRVLKY 15

DB 541 AFRALIQITIFLQY 555

Search completed: February 7, 2000, 18:04:35
Job time: 22201 sec

OS Candida maltosa (Yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Candidaceae; Candida.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=ATCC 20184;
 RX MEDLINE; 88124223.
 RT "Complete nucleotide sequence of the peroxisomal acyl CoA oxidase
 from the alkane-utilizing yeast *Candida maltosa*.";
 RL Nucleic Acids Res. 16:363-366(1988).
 CC -|- CATALYTIC ACTIVITY: ACYL-CoA + O(2) = TRANS-2,3-DEHYDROACYL-CoA +
 H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
 FROM 8 TO 18).
 CC -|- COFACTOR: FAD.
 CC -|- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
 CC SYSTEM.
 CC -|- SUBUNIT: HOMOOCTAMER.
 CC -|- SUBCELLULAR LOCATION: PEROXISOMAL.
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 DR EMBL; X06721; CAA29901.1; -
 DR PIR; A29441; OXCKPM.
 KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
 FT MULTIGENE FAMILY.
 FT INIT_MET 0 BY SIMILARITY.
 SQ SEQUENCE 708 AA; 78242 MW; D5E344D2 CRC32;
 Query Match 54.3%; Score 38; DB 1; Length 708;
 Best Local Similarity 46.7%; Pred. No. 17;
 Matches 7; Conservative 4; Mismatches 4; Indels 0; Gaps 0;
 YQ 1 AYRLIKVIRIVLKY 15
 DB 306 SYRLARVSTIALRY 320
 RESULT 3
 Y4RI_RHISN STANDARD; PRT; 390 AA.
 AC P55642;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 01-NOV-1997 (Rel. 35, Last annotation update)
 DE HYPOTHETICAL 44.0 KD PROTEIN Y4RI.
 GN Y4RI.
 OS Rhizobium sp. (strain NGR234).
 OC Bacterium; Proteobacteria; alpha subdivision; Rhizobiaceae group;
 OC Rhizobiaceae; Rhizobium.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 97305936.
 RA FRIEBERG C.A., FELLAY R., BAIRDOCH A., BROUGHTON W.J., ROSENTHAL A.,
 RA PERRET X.;
 RT "Molecular basis of symbiosis between *Rhizobium* and legumes.";
 RL Nature 387:394-401(1997).
 CC -|- SIMILARITY: NONE OBVIOUS.
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 CC EMBL; AE00094; AAB91834.1; -
 KW Hypothetical protein; Plasmid.
 SQ SEQUENCE 390 AA; 43978 MW; F0AF4E11 CRC32;
 Query Match 52.9%; Score 37; DB 1; Length 390;
 Best Local Similarity 50.0%; Pred. No. 14;
 Matches 7; Conservative 2; Mismatches 5; Indels 0; Gaps 0;
 YQ 2 YRLIKVIRIVLKY 15
 DB 157 YRELLKIARTLSY 170
 RESULT 4
 YM76_YEAST STANDARD; PRT; 724 AA.
 AC Q05029;
 DT 01-NOV-1997 (Rel. 35, Created)
 DT 01-NOV-1997 (Rel. 35, Last sequence update)
 DT 01-NOV-1997 (Rel. 35, Last annotation update)
 DE HYPOTHETICAL 82.0 KD PROTEIN IN RNAI-RNT1 INTERGENIC REGION.
 GN YMR237W OR YMR959.19.
 OS Saccharomycetes cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Saccharomycetes.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=S288C / AB972;
 RA SKELTON J., CHURCHER C.M., BARRELL B.G., RAJANDREAM M.A., WALSH S.V.;
 RL Submitted (JUN-1995) to the EMBL/GenBank/DBJ databases.
 CC -|- SIMILARITY: TO YEAST CSD3/CHS6 AND YEAST YKR027W.
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 or send an email to license@isb-sib.ch).
 CC or send an email to license@isb-sib.ch.
 CC EMBL; Z49939; CAA90208.1; -
 KW Hypothetical protein.
 SQ SEQUENCE 724 AA; 82048 MW; 6CE22AD2 CRC32;
 Query Match 52.1%; Score 36.5; DB 1; Length 724;
 Best Local Similarity 40.0%; Pred. No. 33;
 Matches 8; Conservative 6; Mismatches 1; Indels 5; Gaps 1;
 YQ 1 AYRLIKVIRI-----VLKY 15
 DB 480 AYRLTEIVQITGWQLKY 499
 RESULT 5
 GTA4_MOUSE STANDARD; PRT; 222 AA.
 ID GTA4_MOUSE
 AC P24472;
 DT 01-MAR-1992 (Rel. 21, Created)
 DT 01-OCT-1993 (Rel. 27, Last sequence update)
 DT 15-DEC-1999 (Rel. 39, Last annotation update)
 DE GLUTATHIONE S-TRANSFERASE 5.7 (EC 2.5.1.18) (GST 5.7) (GST CLASS-
 DE ALPHA) (GST A4-4) (GSTA4-4).
 GN GSTA.
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
 OC Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=LUNG;

```

RX MEDLINE; 93050245.
RA ZIMNIAK P., ECKLES M.A., SAXENA M., AWASTHI Y.C.;
RT "A subgroup of class alpha glutathione S-transferases. Cloning of
RL cDNA for mouse lung glutathione S-transferase GST 5.7.";
RN FEBS Lett. 313:173-176(1992).
[2]
RP SEQUENCE OF 106-120 AND 167-184.
RC STRAIN=CD-1; TISSUE=LUNG, AND LIVER;
RX MEDLINE; 91378941.
RA MEDH R.D., SAXENA M., SINGHAL S.S., AHMAD H., AWASTHI Y.C.;
RT "Characterization of a novel glutathione S-transferase isoenzyme from
RL mouse lung and liver having structural similarity to rat glutathione
RN S-transferase 8-8.";
RX Biochem. J. 278:793-799(1991).
[3]
RP X-RAY CRYSTALLOGRAPHY (2.9 ANGSTROMS).
RC TISSUE=LUNG;
RX MEDLINE; 98158566.
RA KRENGEL U., SCHROTER K.H., HOIER H., ARKEMA A., KALK K.H., ZIMNIAK P.,
RA DIJKSTRA B.W.;
RT "Crystal structure of a murine alpha-class glutathione S-transferase
RL involved in cellular defense against oxidative stress.";
RN FEBS Lett. 422:285-290(1998).
[4]
RP X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS).
RC TISSUE=LUNG;
RX MEDLINE; 99438340.
RA XIAO B., SINGH S.P., NANDURI B., AWASTHI Y.C., ZIMNIAK P., JI X.;
RT "Crystal structure of a murine glutathione S-transferase in complex
RL with a glutathione conjugate of 4-hydroxynon-2-enal in one subunit
RN and glutathione in the other: evidence of signaling across the dimer
interface.";
RL Biochemistry 38:11887-11894(1999).
CC -!- FUNCTION: CONJUGATION OF REDUCED GLUTATHIONE TO A WIDE NUMBER
CC OF EXOGENOUS AND ENDOGENOUS HYDROPHOBIC ELECTROPHILES
CC -!- CATALYTIC ACTIVITY: RX + GLUTATHIONE = HX + R-S-GLUTATHIONE.
CC -!- SUBUNIT: HOMODIMER.
CC -!- SUBCELLULAR LOCATION: CYTOPLASMIC.
CC -!- PTM: THE N-TERMINUS IS BLOCKED.
CC -!- MISCELLANEOUS: ON THE BASIS OF IMMUNOLOGICAL AND KINETICS DATA,
CC GST 5.7 IS DISTINCT FROM ALPHA, MU AND PI CLASSES OF GSTS. HOWEVER
CC IT HAS BEEN POSTULATED THAT THIS PROTEIN MAY BE PART OF A DISTINCT
CC SUBGROUP WITHIN THIS ALPHA CLASS.
CC -!- MISCELLANEOUS: THE VARIATIONS WERE FOUND FROM AA SEQUENCING AND
CC IMPLY THERE ARE MULTIPLE FORMS OF THIS PROTEIN. THESE VARIATIONS
CC ARE LIKELY TO BE SEX-LINKED AND TISSUE SPECIFIC.
CC -!- SIMILARITY: BELONGS TO THE GST SUPERFAMILY. ALPHA FAMILY.
-----
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-----
CC EMBL; L06047; AAA37754.1;
CC PIR; S27234; S27234.
CC PDB; 1GUK; 08-APR-98.
CC PDB; 1B48; 29-SEP-99.
CC MGD; MGI:95957; GSTA.
CC PFAM; PF00043; GST; 1.
KW Transferase; Multigene family; Polymorphism; 3D-structure.
FT VARIANT 115 115 K -> P.
FT VARIANT 167 167 V -> G.
FT VARIANT 179 180 PL -> GE.
SQ SEQUENCE 222 AA; 25576 MW; 7158E30C CRC32;

Query Match 51.4%; Score 36; DB 1; Length 222;
Best Local Similarity 60.0%; Pred. No. 13;
Matches 6; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

CC EMBL; L06047; AAA37754.1;
CC PIR; S27234; S27234.
CC PDB; 1GUK; 08-APR-98.
CC PDB; 1B48; 29-SEP-99.
CC MGD; MGI:95957; GSTA.
CC PFAM; PF00043; GST; 1.
KW Transferase; Multigene family; Polymorphism; 3D-structure.
FT VARIANT 115 115 K -> P.
FT VARIANT 167 167 V -> G.
FT VARIANT 179 180 PL -> GE.
SQ SEQUENCE 222 AA; 25576 MW; 7158E30C CRC32;

Query Match 51.4%; Score 36; DB 1; Length 222;
Best Local Similarity 60.0%; Pred. No. 13;
Matches 6; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

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-----
CC EMBL; X75950; CAA33551.1;
CC EMBL; Z28222; CAA82067.1;
CC PIR; S38066; S38066.
CC PIR; S43548; S43548.
CC HSSP; P12351; 1PYC.
CC PROSITE; PS00463; ZN2_CY6_FUNGAL_1; 1.
CC PROSITE; PS50048; ZN2_CY6_FUNGAL_2; 1.
CC PFAM; PF00172; Zn_c1us; 1.
KW Hypothetical protein; Transcription regulation; DNA-binding;
KW Nuclear protein; Zinc; Metal-binding.
FT DNA_BIND 24 52 ZN(2)-CYS(6); FUNGAL-TYPE.
SQ SEQUENCE 705 AA; 82248 MW; 9663DA3B CRC32;

Query Match 51.4%; Score 36; DB 1; Length 705;
Best Local Similarity 46.7%; Pred. No. 40;
Matches 7; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

CC 1 AYRLIKVIRIVLKY 15
CC : : : : :
CC 541 AFRALQIYVIFLQY 555
-----
RESULT 7
SRB8_YEAST
ID SRB8_YEAST STANDARD; PRT; 1427 AA.
AC P25648;
DT 01-MAY-1992 (Rel. 22, Created)
DT 13-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE SUPPRESSOR OF RNA POLYMERASE B SRB8.
GN SRB8 OR YCR081W OR YCR81W.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomyces.
RN [1]

QY 6 IKVIRIVLKY 15
Db 213 VEVVRIVLKF 222
: : : : :
: : : : :
-----
RESULT 6
YKW2_YEAST
ID YKW2_YEAST STANDARD; PRT; 705 AA.
AC P35995;
DT 01-JUN-1994 (Rel. 29, Created)
DT 01-JUN-1994 (Rel. 29, Last sequence update)
DT 01-OCT-1994 (Rel. 30, Last annotation update)
DE PUTATIVE 82.2 KD TRANSCRIPTIONAL REGULATORY PROTEIN IN FRE2 5' REGION.
GN YKL222C.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomyces.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94378726.
RA ALEXANDRAKI D., TZERMIA M.;
RT "Sequencing of a 13.2 kb segment next to the left telomere of yeast
RL chromosome XI revealed five open reading frames and recent
RN recombination events with the right arms of chromosomes III and V.";
RX Yeast 10:S81-S91(1994).
CC -!- SUBCELLULAR LOCATION: NUCLEAR (PROBABLE).
CC -!- SIMILARITY: CONTAINS A ZN(2)-CYS(6), FUNGAL-TYPE BINUCLEAR
CC CLUSTER DOMAIN.
-----
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-----
CC EMBL; X75950; CAA33551.1;
CC EMBL; Z28222; CAA82067.1;
CC PIR; S38066; S38066.
CC PIR; S43548; S43548.
CC HSSP; P12351; 1PYC.
CC PROSITE; PS00463; ZN2_CY6_FUNGAL_1; 1.
CC PROSITE; PS50048; ZN2_CY6_FUNGAL_2; 1.
CC PFAM; PF00172; Zn_c1us; 1.
KW Hypothetical protein; Transcription regulation; DNA-binding;
KW Nuclear protein; Zinc; Metal-binding.
FT DNA_BIND 24 52 ZN(2)-CYS(6); FUNGAL-TYPE.
SQ SEQUENCE 705 AA; 82248 MW; 9663DA3B CRC32;

Query Match 51.4%; Score 36; DB 1; Length 705;
Best Local Similarity 46.7%; Pred. No. 40;
Matches 7; Conservative 4; Mismatches 4; Indels 0; Gaps 0;

CC 1 AYRLIKVIRIVLKY 15
CC : : : : :
CC 541 AFRALQIYVIFLQY 555
-----
RESULT 7
SRB8_YEAST
ID SRB8_YEAST STANDARD; PRT; 1427 AA.
AC P25648;
DT 01-MAY-1992 (Rel. 22, Created)
DT 13-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE SUPPRESSOR OF RNA POLYMERASE B SRB8.
GN SRB8 OR YCR081W OR YCR81W.
OS Saccharomyces cerevisiae (Baker's yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Saccharomycetaceae; Saccharomyces.
RN [1]
```

RP SEQUENCE FROM N.A.
RX MEDLINE: 95293223.
RA HENGARTNER C.J., THOMPSON C.M., ZHANG J., CHAO D.M., LIAO S.M.,
RA KOLSKIE A.J., OKAMURA S., YOUNG R.A.;
RT "Association of an activator with an RNA polymerase II holoenzyme.";
RL Genes Dev. 9:897-910(1995).
RN [2]
RN REVISIONS.
RA GROMADKA R.;
RL Submitted (JAN-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RN SEQUENCE OF 531-1427 FROM N.A.
RA FELDMANN H., MANNAHUT G., VETTER I.;
RL Submitted (MAR-1992) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: COMPONENT OF THE RNA POLYMERASE II HOLOENZYME AND THE
CC MEDIATOR OF ACTIVATION SUBCOMPLEX.
CC -1- SUBCELLULAR LOCATION: NUCLEAR (PROBABLE).
CC -----
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CC -----
DR EMBL: X59720; CAA42268.1; -
DR PIR: S19496; S19496.
DR TRANSFAC: T02152; -
DR SGD: L0003007; SRB8.
KW Nuclear protein.
SQ SEQUENCE 1427 AA; 166859 MW; 6B732E51 CRC32;

Query Match 51.4%; Score 36; DB 1; Length 1427;
Best Local Similarity 77.8%; Pred. No. 81;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 2 YRLIKVIR 10
||| ||| |||
Db 1200 YHLLIKIIR 1208

RESULT 8
RRPL_SENDE
ID RRPL_SENDE STANDARD; PRT; 2048 AA.
AC P06829;
DT 01-JAN-1988 (Rel. 06, Created)
DT 01-JAN-1988 (Rel. 06, Last sequence update)
DT 15-DEC-1998 (Rel. 37, Last annotation update)
DE RNA POLYMERASE BETA SUBUNIT (EC 2.7.7.48) (LARGE STRUCTURAL PROTEIN)
DE (L PROTEIN).
GN L.

OS Sendai virus (strain Enders).
OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Paramyxovirus.
RN [1]

RP SEQUENCE FROM N.A.
RX MEDLINE: 86317720.

RA MORGAN E.M., RAKESTRAW K.M.;
RT "Sequence of the Sendai virus L gene: open reading frames upstream of
RT the main coding region suggest that the gene may be polycistronic.";
RL Virology 154:31-40(1986).

CC -1- FUNCTION: PROBABLE COMPONENT OF THE ACTIVE POLYMERASE. IT MAY
CC FUNCTION IN MRNA SYNTHESIS, CAPPING, METHYLATION AND POLY(A)
CC SYNTHESIS OF NEWLY SYNTHESIZED VIRAL MRNAS. RNA EDITING OF THE P
CC GENE TRANSCRIPT, AND PROTEIN KINASE ACTIVITY.
CC -1- MISCELLANEOUS: THEY ARE FOUND IN CATALYTIC AMOUNTS (APPROXIMATELY
CC 20 TO 30 COPIES PER SENDAI VIRION) IN VIRAL NUCLEOCAPSIDS.
CC -1- SIMILARITY: WITH L PROTEIN OF OTHER PARAMYXOVIRUSES.
CC -----

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CC -----

DR EMBL: M14887; AAA69579.1; -
DR EMBL: D00053; BAA00036.1; -
DR PIR: A24293; ZLNZSE.
DR PFAM: PF00946; Paramyx_RNA_pol; 1.
KW transferase; RNA-directed RNA polymerase.
SQ SEQUENCE 2048 AA; 231623 MW; 7FA48C13 CRC32;

Query Match 51.4%; Score 36; DB 1; Length 2048;
Best Local Similarity 53.8%; Pred. No. 1.2e+02;
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

OY 2 YRLIKVIRIVLK 14
||| ||| ||| |||
Db 2003 YRFLTKIKILMK 2015

RESULT 9

RRPL_SENDS
ID RRPL_SENDS STANDARD; PRT; 2228 AA.

AC P27566;
DT 01-AUG-1992 (Rel. 23, Created)
DT 15-JUL-1999 (Rel. 38, Last sequence update)
DT 15-JUL-1999 (Rel. 38, Last annotation update)
DE RNA POLYMERASE BETA SUBUNIT (EC 2.7.7.48) (LARGE STRUCTURAL PROTEIN)
DE (L PROTEIN).
GN L.

OS Sendai virus (strain Z / host mutants).

OC Viruses; ssRNA negative-strand viruses; Mononegavirales;
OC Paramyxoviridae; Paramyxovirinae; Paramyxovirus.
RN [1]

RP SEQUENCE FROM N.A.
RX STRAIN-MUTANTS TS-F1 AND F1-R;
RX MEDLINE: 90266486

RA MIDDLETON Y., TASHIRO M., THAI T., OH J., SEYMOUR J., PRITZER E.,
RA KLENK H.D., ROTT R., SETO J.T.;

RT "Nucleotide sequence analyses of the genes encoding the HN, M, NP, P,
RT and L proteins of two host range mutants of Sendai virus.";
RL Virology 176:656-657(1990).
RN [2]

RP REVISIONS TO 581 AND 971.
RA MIDDLETON Y.;

RL Submitted (OCT-1998) to the EMBL/GenBank/DBJ databases.
RN [3]

RP SEQUENCE FROM N.A.
RX STRAIN-MUTANT F1-R / T-5 REVERTANT;
RX MEDLINE: 91335752.

RA TASHIRO M., JAMES I., KARRI S., WAHN K., TOBITA K., KLENK H.D.,
RA ROTT R., SETO J.T.;

RT "Pneumotropic revertants derived from a pantropic mutant, F1-R, of
RT Sendai virus.";

RL Virology 184:227-234(1991).

CC -1- FUNCTION: PROBABLE COMPONENT OF THE ACTIVE POLYMERASE. IT MAY
CC FUNCTION IN MRNA SYNTHESIS, CAPPING, METHYLATION AND POLY(A)
CC SYNTHESIS OF NEWLY SYNTHESIZED VIRAL MRNAS. RNA EDITING OF THE P
CC GENE TRANSCRIPT, AND PROTEIN KINASE ACTIVITY.
CC -1- MISCELLANEOUS: THEY ARE FOUND IN CATALYTIC AMOUNTS (APPROXIMATELY
CC 20 TO 30 COPIES PER SENDAI VIRION) IN VIRAL NUCLEOCAPSIDS.
CC -1- SIMILARITY: WITH L PROTEIN OF OTHER PARAMYXOVIRUSES.
CC -----

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P06447;
01-JAN-1988 (Rel. 06, Created)
01-JAN-1988 (Rel. 06, Last sequence update)
15-DEC-1998 (Rel. 37, Last annotation update)
RNA POLYMERASE BETA SUBUNIT (EC 2.7.7.48) (LARGE STRUCTURAL PROTEIN)
(L PROTEIN).
L.
Sendai virus (strain Z).
Viruses; ssRNA negative-strand viruses; Mononegavirales;
Paramyxoviridae; Paramyxovirinae; Paramyxovirus.
[1]
RX MEDLINE; 86148492.
RA SHIODA T., IWASAKI K., SHIBUTA H.;
RT "Determination of the complete nucleotide sequence of the Sendai
RT virus genome RNA and the predicted amino acid sequences of the F, HN
RT and L proteins.";
RL Nucleic Acids Res. 14:1545-1563(1986).
CC -1- FUNCTION: PROBABLE COMPONENT OF THE ACTIVE POLYMERASE. IT MAY
CC FUNCTION IN RNA SYNTHESIS, CAPPING, METHYLATION AND POLY(A)
CC SYNTHESIS OF NEWLY SYNTHESIZED VIRAL MRNAs, RNA EDITING OF THE P
CC GENE TRANSCRIPT, AND PROTEIN KINASE ACTIVITY.
CC -1- MISCELLANEOUS: THEY ARE FOUND IN CATALYTIC AMOUNTS (APPROXIMATELY
CC 20 TO 30 COPIES PER SENDAI VIRION) IN VIRAL NUCLEOCAPSIDS.
CC -1- SIMILARITY: WITH L PROTEIN OF OTHER PARAMYXOVIRUSES.

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DR EMBL: X03614; CAA27273.1; -
DR PIR: A04120; ZLNZSV.
DR PFAM: PF00946; Paramyx_RNA_pol; 1.
KW Transferase: RNA-directed RNA polymerase.
SQ SEQUENCE 228 AA; 252864 MW; DD9798FD CRC32;

Query Match 51.4%; Score 36; DB 1; Length 2228;
Best Local Similarity 53.8%; Pred. No. 1.3e+02;
Matches 7; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 YRLLIKVIRVLK 14
||| |||:|:
DB 2183 YRFLTEIKILMK 2195

RESULT 12
ATPB_MYCLE
ID ATPB_MYCLE STANDARD; PRT; 485 AA.
AC F45823;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 01-NOV-1995 (Rel. 32, Last annotation update)
DE ATP SYNTHASE BETA CHAIN (EC 3.6.1.34).
GN ATPD.
OS Mycobacterium leprae.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RA SMITH D.R., ROBISON K.;
RL Submitted (SEP-1994) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON
CC GRADIENT ACROSS THE MEMBRANE. THE BETA CHAIN IS THE CATALYTIC
CC SUBUNIT.
CC -1- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC
CC CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
CC HAS THREE MAIN SUBUNITS: A, B AND C.
CC

CC -1- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
CC
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CC
CC ENBL; U15186; AAA63108.1; -
DR HSP; P07677; ISKY.
DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
DR PFAM; PF00006; ATP-synt_ab; 1.
DR PFAM; PF00306; ATP-synt_ab.C; 1.
KW Hydrolase; ATP synthesis; CF(1); ATP-binding;
KW Hydrogen ion transport.
FT NP_BIND 170 177 ATP (POTENTIAL).
SQ SEQUENCE 485 AA; 53034 MW; 07216783 CRC32;

Query Match 50.0%; Score 35; DB 1; Length 485;
Best Local Similarity 50.0%; Pred. No. 42;
Matches 7; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 2 YRLIKVIRIVLKY 15
Db 376 YRVAQEVIRILQRY 389
|||:||||:|

RESULT 13
ATPB_MYCTU STANDARD; PRT; 486 AA.
AC Q10593;
DT 01-OCT-1996 (Rel. 34, Created)
DT 01-OCT-1996 (Rel. 34, Last sequence update)
DT 15-DEC-1999 (Rel. 39, Last annotation update)
DE ATP SYNTHASE BETA CHAIN (EC 3.6.1.34).
GN ATPD OR RV1310 OR MTCY373.30.
OS Mycobacterium tuberculosis.
OC Bacteria; Firmicutes; Actinobacteria; Actinobacteridae;
OC Actinomycetales; Corynebacterineae; Mycobacteriaceae; Mycobacterium.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H37RV;
RX MEDLINE; 98295987.
RA COLE S.T., BROSCHE R., PARKHILL J., GARNIER T., CHURCHER C., HARRIS D.,
RA GORDON S.V., EIGLMEIER K., GAS S., BARRY C.E. III, TEKAIA F.,
RA BADCOCK K., BASHAM D., BROWN D., CHILLINGWORTH T., CONNOR R.,
RA DAVIES R., DEVLIN K., FELTWELL T., GENTLES S., HAMLIN N., HOLROYD S.,
RA HORNSBY T., JAGELS K., KROGH A., MCLEAN J., MOULE S., MURPHY L.,
RA OLIVER S., OSBORNE J., QUAIL M.A., RAJANDREAM M.A., ROGERS J.,
RA RUTTER S., SEEGER K., SKELTON S., SQUARES S., SOARES R., SULSTON J.E.,
RA TAYLOR K., WHITEHEAD S., BARRELL B.G.;
RT "Deciphering the biology of Mycobacterium tuberculosis from the
RT complete genome sequence."
RL Nature 393:537-544(1998).
CC -1- FUNCTION: PRODUCES ATP FROM ADP IN THE PRESENCE OF A PROTON
CC GRADIENT ACROSS THE MEMBRANE. THE BETA CHAIN IS THE CATALYTIC
CC SUBUNIT.
CC -1- SUBUNIT: F-TYPE ATPASES HAVE 2 COMPONENTS, CF(1) - THE CATALYTIC
CC CORE - AND CF(0) - THE MEMBRANE PROTON CHANNEL. CF(1) HAS FIVE
CC SUBUNITS: ALPHA(3), BETA(3), GAMMA(1), DELTA(1), EPSILON(1). CF(0)
CC HAS THREE MAIN SUBUNITS: A, B AND C (BY SIMILARITY).
CC -1- SIMILARITY: BELONGS TO THE ATPASE ALPHA/BETA CHAINS FAMILY.
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CC ENBL; Z73419; CAA97743.1; -
DR HSP; P07677; ISKY.
DR PROSITE; PS00152; ATPASE_ALPHA_BETA; 1.
DR PFAM; PF00006; ATP-synt_ab; 1.
DR PFAM; PF00306; ATP-synt_ab.C; 1.
KW Hydrolase; ATP synthesis; CF(1); ATP-binding;
KW Hydrogen ion transport.
FT NP_BIND 171 178 ATP (POTENTIAL).
SQ SEQUENCE 486 AA; 53094 MW; A8001B2F CRC32;

Query Match 50.0%; Score 35; DB 1; Length 486;
Best Local Similarity 50.0%; Pred. No. 42;
Matches 7; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

Qy 2 YRLIKVIRIVLKY 15
Db 377 YRVAQEVIRILQRY 390
|||:||||:|

RESULT 14
CAO3_CANTR STANDARD; PRT; 502 AA.
AC P11355;
DT 01-JUL-1989 (Rel. 11, Created)
DT 01-JUL-1989 (Rel. 11, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE ACYL-COENZYME A OXIDASE POX4-2 (EC 1.3.3.6) (ACYL-COA OXIDASE)
DE (FRAGMENT).
GN POX4-2.
OS Candida tropicalis (Yeast).
OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
OC Candidaceae; Candida.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 87280361.
RA SMALL G.M., LAZAROW P.B.;
RT "Import of the carboxy-terminal portion of acyl-CoA oxidase into
RT peroxisomes of Candida tropicalis."
RL J. Cell Biol. 105:247-250(1987).
CC -1- CATALYTIC ACTIVITY: ACYL-COA + O(2) - TRANS-2,3-DEHYDROACYL-COA +
CC H(2)O(2) (ACTS ON COA DERIVATIVES OF FATTY ACIDS WITH CHAIN LENGTH
CC FROM 8 TO 18).
CC -1- COFACTOR: FAD.
CC -1- PATHWAY: INITIAL STEP OF THE PEROXISOMAL FATTY ACID BETA-OXIDATION
CC SYSTEM.
CC -1- SUBUNIT: HOMOOCTAMER.
CC -1- SUBCELLULAR LOCATION: PEROXISOMAL.
DR PIR; A28584; OXCKAX.
KW Oxidoreductase; Fatty acid metabolism; Peroxisome; Flavoprotein; FAD;
KW Multigene family.
FT NON_TER 1
FT SEQUENCE 502 AA; 55528 MW; EACE80C4 CRC32;

Query Match 50.0%; Score 35; DB 1; Length 502;
Best Local Similarity 40.0%; Pred. No. 43;
Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AYRLIKVIRIVLKY 15
Db 100 SYRLARMSTIALRY 114
|||:|:|:|

RESULT 15
CAO2_CANTR STANDARD; PRT; 708 AA.
AC P06598;
DT 01-JAN-1988 (Rel. 06, Created)
DT 01-NOV-1988 (Rel. 09, Last sequence update)
DT 01-NOV-1997 (Rel. 35, Last annotation update)
DE ACYL-COENZYME A OXIDASE II (EC 1.3.3.6) (ACYL-COA OXIDASE) (PXP-4).
DE

Db 306 SYRMLARMSTIALRY 320

Search completed: February 8, 2000, 01:25:56
Job time: 1556 sec

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Query Match          50.0%; Score 35; DB 1; Length 708;
Best Local Similarity 40.0%; Pred. No. 61;
Matches 6; Conservative 5; Mismatches 4; Indels 0; Gaps 0;

QY      1  AYRLIKIVIRVLKY 15
      :||| : : | | :|

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GenCore version 4.5
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OM protein - protein search, using sw model

Run on: February 8, 2000, 13:17:53 ; Search time 176.54 Seconds
(without alignments)
5.891 Million cell updates/sec

Title: US-08-653-294-21
Perfect score: 70
Sequence: 1 AYRLIKVIRIVLKY 15

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 225878 seqs, 69334122 residues

Total number of hits satisfying chosen parameters: 225878

Minimum DB seq length: 0
Maximum DB seq length: 1000000

Post-processing: Minimum Match 0%
Listing first 45 summaries

Database :

SPTREMBL_12:*

- 1: sp_archaea:*
- 2: sp_bacteria:*
- 3: sp_fungi:*
- 4: sp_human:*
- 5: sp_invertebrate:*
- 6: sp_mammal:*
- 7: sp_mhc:*
- 8: sp_organelle:*
- 9: sp_phage:*
- 10: sp_plant:*
- 11: sp_rodent:*
- 12: sp_virus:*
- 13: sp_vertebrate:*
- 14: sp_unclassified:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	40	57.1	2810	5 Q20456	Q20456 caenorhabdi
2	39.5	56.4	746	3 Q08754	Q08754 saccharomyc
3	38	54.3	771	5 Q20026	Q20026 caenorhabdi
4	37	52.9	98	2 Q45637	Q45637 bacillus sp
5	37	52.9	1696	5 Q21436	Q21436 caenorhabdi
6	36	51.4	100	2 Q69234	Q69234 bacillus ce
7	36	51.4	131	2 Q66643	Q66643 aquifex aeo
8	36	51.4	141	4 Q75379	Q75379 homo sapien
9	36	51.4	186	5 Q96287	Q96287 plasmodium
10	36	51.4	187	2 Q84836	Q84836 chlanydia t
11	36	51.4	282	1 Q30154	Q30154 archaeoglob
12	36	51.4	284	2 Q50343	Q50343 lactobacill
13	36	51.4	325	2 Q929X8	Q929X8 frateuria s
14	36	51.4	381	2 Q86996	Q86996 acinetobact
15	36	51.4	492	3 Q08966	Q08966 saccharomyc
16	36	51.4	1980	12 Q84185	Q84185 human parai
17	36	51.4	2018	5 Q20487	Q20487 caenorhabdi
18	36	51.4	2223	12 Q9WF25	Q9WF25 human parai
19	36	51.4	2228	12 Q98705	Q98705 sendai viru
20	35.5	50.7	1036	10 Q49323	Q49323 arabidopsis

21	35	50.0	83	12	Q89074	Q89074 variola vir
22	35	50.0	175	5	Q17265	Q17265 brugia paha
23	35	50.0	240	2	Q68993	Q68993 chlorobium
24	35	50.0	341	1	Q9VDV4	Q9VDV4 aeropyrum p
25	35	50.0	555	5	Q76770	Q76770 dictyosteli
26	35	50.0	791	10	Q80962	Q80962 arabidopsis
27	35	50.0	1687	11	Q9WUF6	Q9WUF6 mus musculu
28	35	50.0	1740	5	Q23891	Q23891 dictyosteli
29	35	50.0	1740	8	Q9XPI9	Q9XPI9 dictyosteli
30	35	50.0	1782	13	Q42180	Q42180 fugu rubrip
31	35	50.0	1784	4	Q75275	Q75275 homo sapien
32	35	50.0	1786	5	Q17344	Q17344 caenorhabdi
33	35	50.0	1809	5	Q17487	Q17487 caenorhabdi
34	35	50.0	1814	11	Q61037	Q61037 mus musculu
35	35	50.0	1815	5	Q17488	Q17488 caenorhabdi
36	35	50.0	1827	5	Q97275	Q97275 plasmodium
37	35	50.0	1867	5	Q17486	Q17486 caenorhabdi
38	35	50.0	2039	5	Q17489	Q17489 caenorhabdi
39	35	50.0	2228	12	Q55528	Q55528 sendai viru
40	35	50.0	2228	12	Q55530	Q55530 sendai viru
41	35	50.0	3973	5	Q96204	Q96204 plasmodium
42	35	50.0	6994	5	Q17343	Q17343 caenorhabdi
43	34	48.6	180	1	Q9YC59	Q9YC59 aeropyrum p
44	34	48.6	185	2	Q9Z973	Q9Z973 chlanydia p
45	34	48.6	208	8	Q21527	Q21527 clethrionom

ALIGNMENTS

RESULT 1

Q20456 ID Q20456 PRELIMINARY; PRT; 2810 AA.
AC Q20456;
DT 01-NOV-1996 (TREMREL. 01, Created)
DT 01-NOV-1998 (TREMREL. 08, Last sequence update)
DT 01-MAY-1999 (TREMREL. 10, Last annotation update)
DE HUM-4 PROTEIN.
GN HUM-4.
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RA COTTAGE A.;
RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M.,
RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
RA GARDNER A., GREEN P., HAWKINS J., HILLIER L., JIER M., JOHNSON L.,
RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P.,
RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
RA PARSONS J., PERCY C., RIKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
RA SWALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
RA THIERY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans.";
RL Nature 368:32-38(1994).
DR EMBL; Z66563; CAA91469.1; -
DR PFAM; PF00612; IQ; 2.
DR PFAM; PF00063; myosin_head; 4.
DR PFAM; PF00784; MYTH4; 2.
SQ SEQUENCE 2810 AA; 323526 MW; 62742B6C CRC32;

Query Match 57.1%; Score 40; DB 5; Length 2810;
Best Local Similarity 35.7%; Pred. No. 1.6e+02;
Matches 5; Conservative 7; Mismatches 2; Indels 0; Gaps 0;

QY 2 YRLIKVIRIVLKY 15
 ID 1102 FRLSVEIFKLILAY 1115

RESULT 2
 Q08754 PRELIMINARY; PRT; 746 AA.
 AC Q08754;
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-NOV-1996 (TREMBLrel. 01, Last annotation update)
 DE ORF YOR299W.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Saccharomycetes.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA CZEPLUCH C., JAUNIAUX J.C., KORDS E., POIREY R., PUJOL A.,
 RA TOBIASCH E.;
 RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA MIPS;
 RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
 DR EMBL; 275207; CAA99528.1;
 SQ SEQUENCE 746 AA; 84829 MW; 01352AC2 CRC32;

Query Match 56.4%; Score 39.5; DB 3; Length 746;
 Best Local Similarity 45.0%; Pred. NO. 56;
 Matches 9; Conservative 5; Mismatches 1; Indels 5; Gaps 1;

QY 1 AYELIKVIRI-----VLKY 15
 ID 502 AYRLTEIVOITGWENLLKY 521

RESULT 3
 Q20026 PRELIMINARY; PRT; 771 AA.
 AC Q20026;
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
 DE CODED FOR BY C. ELEGANS CDNA CEESB82F.
 GN F35C8.7.
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
 OC Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE; 94150718.
 RA WILSON R., AINSKOUGH R., ANDERSON K., BAYNES C., BERKS M.,
 RA BONFIELD J., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A.,
 RA CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L.,
 RA GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L.,
 RA JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATRILLE P.,
 RA LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M.,
 RA PARSONS J., PERCY C., RIFKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R.,
 RA SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J.,
 RA THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R.,
 RA WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 elegans.";
 RL Nature 368:32-38(1994).
 RN [2]
 RP SEQUENCE FROM N.A.
 RA WU X.;
 RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RA WATERSTON R.;

RL Submitted (NOV-1995) to the EMBL/GenBank/DBJ databases.
 DR EMBL; U40941; AAA81709.1;
 SQ SEQUENCE 771 AA; 87310 MW; 5AE2EE3F CRC32;

Query Match 54.3%; Score 38; DB 5; Length 771;
 Best Local Similarity 45.5%; Pred. NO. 1.1e+02;
 Matches 5; Conservative 6; Mismatches 0; Indels 0; Gaps 0;

QY 5 LIKIVIRIVLKY 15
 DB 578 IVKIIRVLELEY 588

RESULT 4
 Q45637 PRELIMINARY; PRT; 98 AA.
 ID Q45637;
 AC Q45637;
 DT 01-NOV-1996 (TREMBLrel. 01, Created)
 DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
 DT 01-NOV-1999 (TREMBLrel. 12, Last annotation update)
 DE HYPOTHETICAL 10.9 KD PROTEIN.
 OS Bacillus sp., Bacillus megaterium, and Exiguobacterium sp.
 OC Bacteria; Firmicutes; Bacillus/Clostridium group;
 OC Bacillus/Staphylococcus group; Bacillus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-RC507;
 RX MEDLINE; 89133092.
 RA WANG Y., MOORE M., LEVINSON H.S., SILVER S., WALSH C., MAHLER I.;
 RT "Nucleotide sequence of a chromosomal mercury resistance determinant
 from a Bacillus sp. with broad-spectrum mercury resistance.";
 RL J. Bacteriol. 171:83-92(1989).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-MK64-1;
 RA MINAKHIN L.S.;
 RL Submitted (DEC-1996) to the EMBL/GenBank/DBJ databases.
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN-MK64-1, TC38-2B;
 RX MEDLINE; 98195721.
 RA BOGDANOVA E.S., BASS I.A., MINAKHIN L.S., PETROVA M.A., MINDLIN S.Z.,
 RA VOLODIN A.A., KALYAIEVA E.S., TIEDGE G.M., HOBMAN J.L., BROWN N.L.,
 RA NIKIFIROV V.G.;
 RT "Horizontal spread of mer operons among gram-positive bacteria in
 natural environments.";
 RL Microbiology 144:609-620(1998).
 RN [4]
 RP SEQUENCE FROM N.A.
 RC STRAIN-TC38-2B;
 RA MINAKHIN L.;
 RL Submitted (JUL-1996) to the EMBL/GenBank/DBJ databases.
 RN [5]
 RP SEQUENCE FROM N.A.
 RC STRAIN-TC38-2B;
 RA MINAKHIN L.;
 RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AF138877; AAA83975.1;
 DR EMBL; Y09907; CAA71043.1;
 DR EMBL; X9907; CAA71043.1;
 DR EMBL; X99457; CAA67820.1;
 KW Hypothetical protein; Plasmid.
 SQ SEQUENCE 98 AA; 10900 MW; F1D67BC1 CRC32;

Query Match 52.9%; Score 37; DB 2; Length 98;
 Best Local Similarity 58.3%; Pred. NO. 22;
 Matches 7; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 YRLIKVIRIVL 13
 DB 46 YRLFSIVTIVL 57

RESULT 5
Q21436 PRELIMINARY; PRT; 1696 AA.
AC Q21436;
DT 01-NOV-1996 (TReMBLrel. 01, Created)
DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
DT 01-JAN-1999 (TReMBLrel. 09, Last annotation update)
DE K12D12.2 PROTEIN.
GN K12D12.2
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Secerententea; Rhabditia; Rhabditida; Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
RN [1]
RP SEQUENCE FROM N.A.
RA COLES L.;
RL Submitted (APR-1995) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RX MEDLINE; 94150718.
RA WILSON R., AINSCOUGH R., ANDERSON K., BAYNES C., BERKS M., BURTON J., CONNELL M., COPSEY T., COOPER J., COULSON A., CRAXTON M., DEAR S., DU Z., DURBIN R., FAVELLO A., FULTON L., GARDNER A., GREEN P., HAWKINS T., HILLIER L., JIER M., JOHNSTON L., JONES M., KERSHAW J., KIRSTEN J., LAISTER N., LATREILLE P., LIGHTNING J., LLOYD C., MCMURRAY A., MORTIMORE B., O'CALLAGHAN M., PARSONS J., PERCY C., RIKEN L., ROOPRA A., SAUNDERS D., SHOWNKEEN R., SMALDON N., SMITH A., SONNHAMMER E., STADEN R., SULSTON J., THIERRY-MIEG J., THOMAS K., VAUDIN M., VAUGHAN K., WATERSTON R., WATSON A., WEINSTOCK L., WILKINSON-SPROAT J., WOHLDMAN P.;
RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C. elegans.";
RL Nature 368:32-38(1994).
DR EMBL; 249069; CAA88864.1; -;
SQ SEQUENCE 1696 AA; 191324 MW; 6EDE7750 CRC32;

Query Match 52.9%; Score 37; DB 5; Length 1696;
Best Local Similarity 53.3%; Pred. No. 3.3e+02;
Matches 8; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

QY 1 AYRLIKVIRIVLKY 15
:|||||:|:|:
DB 121 AHRLLIATMRQMLKW 135

RESULT 6
O69234 PRELIMINARY; PRT; 100 AA.
AC O69234;
DT 01-AUG-1998 (TReMBLrel. 07, Created)
DT 01-AUG-1998 (TReMBLrel. 07, Last sequence update)
DT 01-JAN-1999 (TReMBLrel. 09, Last annotation update)
DE TRANSPORT PROTEIN.
OS Bacillus cereus.
OG Plasmid pKUH302, and plasmid pKUH301.
OC Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Bacillus.
RN [1]
RP SEQUENCE FROM N.A.
RC PLASMID-PKUH302; TRANSPONON-TN1546-LIKE, AND TN3-TYPE;
RA MINAKHIN L.S.;
RL Submitted (OCT-1996) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RC PLASMID-PKUH301; TRANSPONON-TN1546-LIKE, AND TN3-TYPE;
RX MEDLINE; 98195721.
RA BOGDANOVA E.S., BASS I.A., MINAKHIN L.S., PETROVA M.A., MINDLIN S.Z., RA VOLODIN A.A., KALVAEVA E.S., TIEDGE G.M., HOBMAN J.L., BROWN N.L., RA NIKIFIROV V.G.;
RT "Horizontal spread of mer operons among gram-positive bacteria in natural environments.";
RL Microbiology 144:609-620(1998).

DR EMBL; Y09024; CAA70222.1; -;
DR EMBL; Y09027; CAA70244.1; -;
KW Plasmid.
SQ SEQUENCE 100 AA; 11126 MW; 381D4576 CRC32;
Query Match 51.4%; Score 36; DB 2; Length 100;
Best Local Similarity 50.0%; Pred. No. 34;
Matches 6; Conservative 3; Mismatches 3; Indels 0; Gaps 0;

QY 2 YRLIKVIRIVL 13
|||||:|:|:
DB 46 YRLFSIVTIL 57

RESULT 7
O66643 PRELIMINARY; PRT; 131 AA.
AC O66643;
DT 01-AUG-1998 (TReMBLrel. 07, Created)
DT 01-AUG-1998 (TReMBLrel. 07, Last sequence update)
DT 01-NOV-1998 (TReMBLrel. 08, Last annotation update)
DE HYPOTHETICAL 15.4 KD PROTEIN.
GN AQ_293.
OS Aquifex aeolicus.
OC Bacteria; Aquificales; Aquificaceae; Aquifex.
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-VF5;
RX MEDLINE; 98196666.
RA DECKERT G., WARREN P.V., GAASTERLAND T., YOUNG W.G., LENOX A.L., GRAHAM D.E., OVERBECK R., SNEAD M.A., KELLER M., AUJAY M., HUBER R., RA FELDMAN R.A., SHORT J.M., OLSON G.J., SWANSON R.V.;
RT "The complete genome of the hyperthermophilic bacterium Aquifex aeolicus.";
RL Nature 392:353-358(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-VF5;
RA DECKERT G., WARREN P.V., GAASTERLAND T., YOUNG W.G., LENOX A.L., GRAHAM D.E., OVERBECK R., SNEAD M.A., KELLER M., AUJAY M., HUBER R., RA FELDMAN R.A., SHORT J.M., OLSON G.J., SWANSON R.V.;
RL Submitted (JUL-1997) to the EMBL/GenBank/DBJ databases.
DR EMBL; AE000682; AAC06601.1; -;
KW Hypothetical protein.
SQ SEQUENCE 131 AA; 15393 MW; E36D7DD3 CRC32;

Query Match 51.4%; Score 36; DB 2; Length 131;
Best Local Similarity 28.6%; Pred. No. 44;
Matches 4; Conservative 8; Mismatches 2; Indels 0; Gaps 0;

QY 2 YRLIKVIRIVLKY 15
:|||||:|:|:
DB 46 HKILVKLVNVIILY 59

RESULT 8
O75379 PRELIMINARY; PRT; 141 AA.
AC O75379;
DT 01-NOV-1998 (TReMBLrel. 08, Created)
DT 01-NOV-1998 (TReMBLrel. 08, Last sequence update)
DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
DE VAMP 4.
GN VAMP4.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia; OC Eutheria; Primates; Catarrhini; Hominidae; Homo.
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE; 98221163.
RA ADVANI R.J., BAE H.R., BOCK J.B., CHAO D.S., DOUNG Y.C., PREKERIS R.,

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RA YOO J.S., SCHELLER R.H.;
RT "Seven novel mammalian SNARE proteins localize to distinct membrane
RL compartments.";
RL J. Biol. Chem. 273:10317-10324(1998).
DR EMBL: AF044310; AAC24032.1; -.
DR PFAM: PF00957; synaptobrevin; 1.
DR PRINTS: PR00219; SYNAPTOBREVN.
SQ SEQUENCE 141 AA; 16366 MW; 3C737E45 CRC32;

Query Match 51.4%; Score 36; DB 4; Length 141;
Best Local Similarity 53.3%; Pred. No. 47;
Matches 8; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AYRLIKVIRIVLKY 15
| | | | |
Db 125 AILLVILVIMKY 139

RESULT 9
O96287 PRELIMINARY; PRT; 186 AA.
AC O96287;
DT 01-MAY-1999 (Tremblrel. 10, Created)
DT 01-MAY-1999 (Tremblrel. 10, Last sequence update)
DT 01-MAY-1999 (Tremblrel. 10, Last annotation update)
DE PREDICTED INTEGRAL MEMBRANE PROTEIN.
GN PF09085C.
OS Plasmodium falciparum.
OC Eukaryota; Alveolata; Apicomplexa; Haemosporidia; Plasmodium.
[1]
RN SEQUENCE FROM N.A.
RX MEDLINE; 99021743.
RA GARDNER M.J., TETTELIN H., CARUCCI D.J., CUMMINGS L.M., ARAVIND L.,
RA KOONIN E.V., SHALLOM S., MASON T., YU K., FUJII C., PEDERSON J.,
RA SHEN K., JING J., ASTON C., LAI Z., SCHWARTZ D.C., PERTEA M.,
RA SALZBERG S., ZHOU L., SUTTON G.G., CLAYTON R., WHITE O., SMITH H.O.,
RA FRASER C.M., ADAMS M.D., VENTER J.C., HOFFMAN S.L.;
RT "Chromosome 2 sequence of the human malaria parasite Plasmodium
falciparum.";
RL Science 282:1126-1132(1998).
DR EMBL: AE001431; AAC71985.1; -.
SQ SEQUENCE 186 AA; 21961 MW; B165839F CRC32;

Query Match 51.4%; Score 36; DB 5; Length 186;
Best Local Similarity 53.3%; Pred. No. 61;
Matches 8; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

Qy 1 AYRLIKVIRIVLKY 15
| | | | |
Db 145 AFILLIFIVHARY 159

RESULT 10
O84836 PRELIMINARY; PRT; 187 AA.
AC O84836;
DT 01-NOV-1998 (Tremblrel. 08, Created)
DT 01-NOV-1998 (Tremblrel. 08, Last sequence update)
DT 01-NOV-1998 (Tremblrel. 08, Last annotation update)
DE RRNA METHYLASE (POSSIBLE).
GN YGGH.
OS Chlamydia trachomatis.
OC Bacteria; Chlamydiales; Chlamydiaceae; Chlamydia.
[1]
RN SEQUENCE FROM N.A.
RX STRAIN=D/UW-3/CX;
RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L.,
RA MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V.,
RA DAVIS R.W.;
RT "Genome Sequence of an Obligate Intracellular Pathogen of Humans:
Chlamydia trachomatis.";
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RL Science 0:0-0(1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=D/UW-3/CX;
RA STEPHENS R.S., KALMAN S., LAMMEL C.J., FAN J., MARATHE R., ARAVIND L.,
RA MITCHELL W.P., OLINGER L., TATISOV R.L., ZHAO Q., KOONIN E.V.,
RA DAVIS R.W.;
RL Submitted (MAY-1998) to the EMBL/GenBank/DBJ databases.
DR EMBL: AE001356; AAC68426.1; -.
KW Methyltransferase.
SQ SEQUENCE 187 AA; 21768 MW; 9AF9D1A5 CRC32;

Query Match 51.4%; Score 36; DB 2; Length 187;
Best Local Similarity 42.9%; Pred. No. 61;
Matches 6; Conservative 3; Mismatches 5; Indels 0; Gaps 0;

Qy 2 YRLIKVIRIVLKY 15
| | | | |
Db 130 YNVLVNAITVIMKY 143

RESULT 11
O30154 PRELIMINARY; PRT; 282 AA.
AC O30154;
DT 01-JAN-1998 (Tremblrel. 05, Created)
DT 01-JAN-1998 (Tremblrel. 05, Last sequence update)
DT 01-AUG-1998 (Tremblrel. 07, Last annotation update)
DE HYPOTHETICAL 33.0 KD PROTEIN.
GN AF0082.
OS Archaeoglobus fulgidus.
OC Archaea; Euryarchaeota; Archaeoglobales; Archaeoglobaceae;
OC Archaeoglobus.
[1]
RN SEQUENCE FROM N.A.
RX STRAIN=VC-16 / DSM 4304 / ATCC 49558;
RX MEDLINE; 98049343.
RA KLENK H.-P., CLAYTON R.A., TOMB J.-F., WHITE O., NELSON K.E.,
RA RICHARDSON D.L., KERLAVAGE A.R., GRAHAM D.E., KYRPIDES N.C.,
RA FLEISCHMANN R.D., QUACKENBUSH J., LEE N.H., SUTTON G.G., GILL S.,
RA KIRKNESS E.F., DOUGHERTY B.A., MCKENNEY K., ADAMS M.D., LOFTUS B.,
RA PETERSON S., REICH C.I., MCNEIL L.K., BADGER J.H., GLODEK A., ZHOU L.,
RA OVERBECK R., GOCAYNE J.D., WEIDMAN J.F., McDONALD L., UTTERBACK T.,
RA COTTON M.D., SPRIGGS T., ARTIACH P., KAINE B.P., SYKES S.M.,
RA SADOW P.W., D'ANDREA K.P., BOWMAN C., FUJII C., GARLAND S.A.,
RA MASON T.M., OLSEN G.J., FRASER C.M., SMITH H.O., WOESE C.R.,
RA VENTER J.C.;
RT "The complete genome sequence of the hyperthermophilic, sulphate-
reducing archaeon Archaeoglobus fulgidus.";
RL Nature 390:364-370(1997).
DR EMBL: AE001100; AAB91148.1; -.
DR TIGR; AF0082; -.
KW Hypothetical protein.
SQ SEQUENCE 282 AA; 33040 MW; 2712078C CRC32;

Query Match 51.4%; Score 36; DB 1; Length 282;
Best Local Similarity 53.8%; Pred. No. 90;
Matches 7; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

Qy 2 YRLIKVIRIVLKY 14
| | | | |
Db 269 YRHLRLIRITLK 281

RESULT 12
O50343 PRELIMINARY; PRT; 284 AA.
ID O50343
AC O50343
DT 01-JUN-1998 (Tremblrel. 06, Created)
DT 01-JUN-1998 (Tremblrel. 06, Last sequence update)
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DT 01-JUN-1998 (TReMBLrel. 06, Last annotation update)
 DE HYPOTHETICAL 33.6 KD PROTEIN.
 OS Lactobacillus helveticus.
 OG Plasmid pHL1
 OC Bacteria; Firmicutes; Bacillus/Clostridium group; Lactobacillaceae;
 OC Lactobacillus.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-ATCC15009;
 RA THOMPSON K., MCCONVILLE K.J., MCREYNOLDS C., FOLEY S.;
 RL Submitted (DEC-1997) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AJ222725; CAA10961.1; -
 KW Hypothetical protein; Plasmid.
 SQ SEQUENCE 284 AA; 33632 MW; E77494B0 CRC32;

Query Match 51.4%; Score 36; DB 2; Length 284;
 Best Local Similarity 28.6%; Pred. No. 91;
 Matches 4; Conservative 8; Mismatches 2; Indels 0; Gaps 0;

QY 2 YRLIKVIRIVLKY 15
 | : : : : :
 Db 231 YKSVKFLKLLIKY 244

RESULT 13
 Q929X8 PRELIMINARY; PRT; 325 AA.
 AC Q929X8;
 DT 01-MAY-1999 (TReMBLrel. 10, Created)
 DT 01-MAY-1999 (TReMBLrel. 10, Last sequence update)
 DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
 DE ELECTRON TRANSFER PROTEIN.
 OS Frateuria sp. ANA-18.
 OC Bacteria; Proteobacteria; gamma subdivision; Frateuria.
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-ANA-18;
 RX MEDLINE; 99132292.
 RA MURAKAMI S., TAKASHIMA A., TAKEMOTO J., TAKENAKA S., SHINKE R.,
 RA AOKI K.;
 RT "Cloning and sequence analysis of two catechol-degrading gene clusters
 from the aniline-assimilating bacterium Frateuria species ANA-18.";
 RL Gene 226:189-198(1999).
 DR EMBL; AB009373; BAA75213.1; -
 DR HSP; P33164; 2PIA.
 DR PROSITE; PS00197; 2FE2S_FERREDOXIN; 1.
 KW Iron-sulfur.
 SQ SEQUENCE 325 AA; 34987 MW; 6A35946C CRC32;

Query Match 51.4%; Score 36; DB 2; Length 325;
 Best Local Similarity 60.0%; Pred. No. 1e+02;
 Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 1 AYRLIKVIRIVLKY 15
 | : : : : :
 Db 127 AYRLKQEQRFVLHY 141

RESULT 14
 O86996 PRELIMINARY; PRT; 381 AA.
 AC O86996;
 DT 01-NOV-1998 (TReMBLrel. 08, Created)
 DT 01-NOV-1998 (TReMBLrel. 08, Last sequence update)
 DT 01-NOV-1999 (TReMBLrel. 12, Last annotation update)
 DE HYPOTHETICAL 41.4 KD PROTEIN.
 OS Acinetobacter lwofii K24.
 OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonas group;
 OC Moraxellaceae; Acinetobacter.
 RN [1]
 RP SEQUENCE FROM N.A.

RC STRAIN-K24;
 RA KIM S.I., LEEM S.-H., CHOI J.S., CHUNG Y.H., KIM S., PARK Y.-M.,
 RA HA K.-S.;
 RL Submitted (AUG-1998) to the EMBL/GenBank/DBJ databases.
 DR EMBL; U77659; AAC31770.1; -
 DR HSP; P33164; 2PIA.
 DR PROSITE; PS00197; 2FE2S_FERREDOXIN; 1.
 DR PFAM; PF00111; fer2; 1.
 KW Hypothetical protein; Iron-sulfur.
 SQ SEQUENCE 381 AA; 41377 MW; BE9CD1C2 CRC32;

Query Match 51.4%; Score 36; DB 2; Length 381;
 Best Local Similarity 60.0%; Pred. No. 1.2e+02;
 Matches 9; Conservative 1; Mismatches 5; Indels 0; Gaps 0;

QY 1 AYRLIKVIRIVLKY 15
 | : : : : :
 Db 183 AYRLKQEQRFVLHY 197

RESULT 15
 Q08966 PRELIMINARY; PRT; 492 AA.
 AC Q08966;
 DT 01-NOV-1996 (TReMBLrel. 01, Created)
 DT 01-NOV-1996 (TReMBLrel. 01, Last sequence update)
 DT 01-NOV-1996 (TReMBLrel. 01, Last annotation update)
 DE CHROMOSOME XVI READING FRAME ORF YPL219W.
 OS Saccharomyces cerevisiae (Baker's yeast).
 OC Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
 OC Saccharomycetaceae; Saccharomyces.
 RN [1]
 RP SEQUENCE FROM N.A.
 RA RIEGER M., MUELLER-AUER S., SCHAEFER M.;
 RL Submitted (JUN-1996) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA MIPS;
 RL Submitted (MAY-1996) to the EMBL/GenBank/DBJ databases.
 DR EMBL; Z73575; CAA97934.1; -
 SQ SEQUENCE 492 AA; 55430 MW; 6C549DE0 CRC32;

Query Match 51.4%; Score 36; DB 3; Length 492;
 Best Local Similarity 50.0%; Pred. No. 1.5e+02;
 Matches 7; Conservative 3; Mismatches 4; Indels 0; Gaps 0;

QY 1 AYRLIKVIRIVLKY 14
 | : : : : :
 Db 416 AHRIITIRIATK 429

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-DELOP=7.000 -YGAPOP=10.000 -YGAPEXT=0.500 -DELOP=6.000
-DELXT=7.000 -START=1 -MATRIX=blosum62 -TRANS=human40.cdi
-LIST=45 -DOALIGN=200 -THR_SCORE=pct -ALIGN=15 -MODE=LOCAL
-OUTFMT=pfs -NORM=ext -MINLEN=0 -MAXLEN=100000 -USER=US08653294
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Search information block:

Query: US-08-653-294-21

Query length: 15

Database: GenEmbl:*

Database sequences: 821193

Database length: -1518192014

Search time (sec): 10176.920000

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gb_p13:AC016012	+	47.00	107.33	590.34	68676
gb_p13:AB017067	+	46.00	102.26	1.1e+03	83689
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gb_p13:AC010014	+	44.00	91.32	4.6e+03	136702
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DEFINITION Arabidopsis thaliana endoxyloglucan transferase (XTR2) gene,
complete cds.
ACCESSION AF163820
VERSION AF163820.1 GI:5533310
KEYWORDS
SOURCE thale cress.

ORGANISM Arabidopsis thaliana
Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons; core
eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
Arabidopsis.

REFERENCE 1 (bases 1 to 3000)
AUTHORS Akamatsu,T., Hanzawa,Y., Ohtake,Y., Takahashi,T., Nishitani,K. and
Komeda,Y.

TITLE Expression of endoxyloglucan transferase genes in acaulis mutants
of Arabidopsis
JOURNAL Plant Physiol. 121 (3), 715-721 (1999)
REFERENCE 2 (bases 1 to 3000)
AUTHORS Takahashi,T.

TITLE Direct Submission
JOURNAL Submitted (30-JUN-1999) Division of Biological Sciences, Graduate
School of Science, Hokkaido University, N10, W8, Sapporo 060-0810,
Japan

FEATURES

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BASE COUNT 906 a 527 c 516 g 1048 t 3 others
ORIGIN

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Ratio: 4.000 Gaps: 0
Percent Similarity: 80.000 Percent Identity: 66.667

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US-08-653-294-21 x AF163820/rev ...

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ACCESSION   Z81552
VERSION     Z81552.1 GI:3217528
KEYWORDS    HTG.
SOURCE      Caenorhabditis elegans.
ORGANISM    Caenorhabditis elegans.
REFERENCE   1 (bases 1 to 38062)
AUTHORS     Wilson,R., Ainscough,R., Anderson,K., Baynes,C., Berks,M.,
            Bonfield,J., Burton,J., Connell,M., Copsey,T., Cooper,J.,
            Coulson,A., Craxton,M., Dear,S., Du,Z., Durbin,R., Favello,A.,
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TITLE       Nature 368 (6466), 32-38 (1994)
JOURNAL     Nature 368 (6466), 32-38 (1994)
MEDLINE     94150718
REFERENCE   2 (bases 1 to 38062)
AUTHORS     Lennard,N.
TITLE       Direct Submission
JOURNAL     Submitted (06-NOV-1996) Louis, MO 63110, USA. E-mail:
            jes@sanger.ac.uk or rwenematode.wustl.edu
COMMENT     On Jun 13, 1998 this sequence version replaced gi:1665968.
            Coding sequences below are predicted from computer analysis, using
            predictions from Genefinder (P. Green, U. Washington), and other
            available information.
            For a graphical representation of this sequence and its analysis
            see:
            http://webace.sanger.ac.uk/cgi-
            bin/display?db=wormacsc&class=Sequence&object=F56G4
            Current sequence finishing criteria for the C. elegans genome
            sequencing consortium are that all bases are either sequenced
            unambiguously on both strands, or on a single strand with both a
            dye primer and dye terminator reaction, from distinct subclones.
            Exceptions are indicated by an explicit note.
            IMPORTANT: This sequence is NOT necessarily the entire insert of
            the specified clone. It may be shorter because we only sequence
            overlapping sections once, or longer because we arrange for a small
            overlap between neighbouring submissions.
            This sequence is the entire insert of clone F56G4. The true right
            end of clone M04D5 is at 19377 in this sequence. The start of this
            sequence (1..100) overlaps with the end of sequence Z83118.
            The end of this sequence (37961..38062) overlaps with the start of
            sequence AL117201.
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ACCESSION AB017067
VERSION AB017067.1 GI:3510343
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eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
Arabidopsis.
REFERENCE 1 (sites)
AUTHORS Nakamura.Y.
TITLE Structural Analysis of Arabidopsis thaliana Chromosome 5. IX
JOURNAL Unpublished (1998)
REFERENCE 2 (bases 1 to 83689)
AUTHORS Nakamura.Y.
TITLE Direct Submission
JOURNAL Submitted (26-AUG-1998) to the DDBJ/EMBL/GenBank databases.
Yasukazu Nakamura, Kazusa DNA Research Institute, Laboratory of
Gene Structure 2, 1532-3, Yana, Kisarazu, Chiba 292, Japan
(E-mail:ynakamu@kazusa.or.jp, Tel:+81-438-52-3935,
Fax:+81-438-52-3934)
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ACCESSION AF027868
VERSION AF027868.1 GI:2618993
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complement(12919. .13617)

alignment_scores:
Quality: 46.00 Length: 13
Ratio: 3.538 Gaps: 0
Percent Similarity: 100.000 Percent Identity: 69.231

alignment_block:
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Align seg 1/1 to: AF027868 from: 1 to: 87500

1 AlaTyrArgLeuLeuIleYsValIleArgIleValLeu 13

35864 AGCTTTAGATTGCTGCTGAAGTAATAAAAAATTGTACTT 35902

seq_name: gb_pl2:ATF13C5

seq_documentation_block:

LOCUS ATF13C5 119111 bp DNA PLN 23-SEP-1999
DEFINITION Arabidopsis thaliana DNA chromosome 4, BAC clone F13C5 (ESSA
project).
ACCESSION AL021711
VERSION AL021711.2 GI:5738363
KEYWORDS

SOURCE

ORGANISM

thale cress.

Arabidopsis thaliana

Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
euphyllophytes; Spermatophyta; Magnoliophyta; eudicotyledons; core
eudicots; Rosidae; eurosids II; Brassicales; Brassicaceae;
Arabidopsi

REFERENCE

AUTHORS

1 (bases 1 to 119111)

Bevan, M., Pohl, T., Weizenegger, T., Bancroft, I., Mewes, H. W.,
Mayer, K. F. X., Lemcke, K. and Schueller, C.

JOURNAL

AUTHORS

2 (bases 1 to 119111)

EU Arabidopsis sequencing project.
Direct Submission

TITLE

JOURNAL

Submitted (23-SEP-1999) MIPS, at the Max-Planck-Institut fuer
Biochemie, Am Klopferspitz 18a, D-82152 Martinsried, FRG, E-mail:
schueller@mips.biochem.mpg.de, mayer@mips.biochem.mpg.de, Project
Coordinator: Mike Bevan, Molecular Genetics Department, Cambridge
Laboratory, John Innes Centre, Colney Lane, NR4 7UJ Norwich, UK,
E-mail: michael.bevan@bbsrc.ac.uk

COMMENT

On Aug 18, 1999 this sequence version replaced gi:2832611.
Information on performance of analysis and a more detailed
annotation of this entry and other sequences of chromosomes 3, 4
and 5 can be viewed at: <http://www.mips.biochem.mpg.de/proj/thal/>.

FEATURES

source

Location/Qualifiers

1. .119111

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/variety="Columbia"

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/chromosome="4"

/note="positions 1-4494 are not included within BAC clone

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2221. .3267

/gene="F13C5.5"

2221. .3267

/gene="F13C5.5"

/note="weak similarity to MICROTUBULE ASSOCIATED PROTEIN

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EQDAGVKKPARPTGTGGYSREDSVLGHTITKPAHOWEKLKVKLRVKKLADQOR
KSLYKRELNRIGTKENKVRVSPRASEKCRVKAIEDLKKAKQAREHLLLETDGQ
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/number=1

3488. .5125

/gene="F13C5.10"

complement(3488. .5125)

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/number=1

complement(3488. .5125)

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/note="similarity to hypothetical proteins - Arabidopsis

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/codon_start=1

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QIHLGFIKSLGTVDEVTLVNNGYGRSGVFEIARKVLDMPVDRDAVSNLSLAVLE
KGLVDCARALFDEMEERNVESNFMISGYAAGLVKAEKVEFDSMPVRDVSNNAMVT
AYAHGCVINEVLEFNKMDDDSTKPDGFTLVSLVLSACLSLGSQGEVHVYIDKRG
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ORGANISM      Homo sapiens
REFERENCE     Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Mammalia;
              Eutheria; Primates; Catarrhini; Hominidae; Homo.
AUTHORS       Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE         Homo sapiens, clone RP11-19D19
JOURNAL       Unpublished
REFERENCE     2 (bases 1 to 160557)
AUTHORS       Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
              Baldwin,J., Barna,N., Beckerly,R., Boguslavskiy,L., Boukhgalter,B.,
              Brown,A., Castelle,A., Colangelo,M., Collins,S., Collymore,A.,
              Cooke,P., DeArelano,K., Dewar,K., Domino,M., Doneelan,L., Doyle,M.,
              Ferreira,P., Fitzhugh,W., Forrest,C., Funke,R., Gage,D.,
              Galagan,J., Gardyna,S., Grant,G., Hagos,B., Hearford,A., Horton,L.,
              Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
              Lehoczy,J., Lieu,C., Locke,K., Macdonald,P., Marquis,N.,
              McEwan,P., McGurk,A., McKernan,K., McLaughlin,J., Meldrim,J.,
              Morrow,J., Naylor,J., Norman,C.H., O'Connor,P., O'Donnell,P.,
              Peterson,K., Pollara,V., Riley,R., Roy,A., Santos,R., Severy,P.,
              Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
              Testaye,S., Tirrell,A., Vassiliev,H., Vo,A., Wheeler,J., Wu,X.,
              Wyman,D., Ye,W.J., Zimmer,A. and Zody,M.
TITLE         Direct Submission
JOURNAL       Submitted (10-OCT-1999) Whitehead Institute/MIT Center for Genome
              Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT       On Nov 30, 1999 this sequence version replaced gi:6018128.
              All repeats were identified using RepeatMasker:
              Smit, A.F.A. & Green, P. (1996-1997)
              http://ftp.genome.washington.edu/RM/RepeatMasker.html
              ----- Genome Center
              Center: Whitehead Institute/ MIT Center for Genome Research
              Center code: WIBR
              Web site: http://www-seq.wi.mit.edu
              Contact: sequence_submission@genome.wi.mit.edu
              ----- Project Information
              Center project name: L3721
              Center clone name: 19_D_19
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              * NOTE: This is a 'working draft' sequence. It currently
              * consists of 33 contigs. The true order of the pieces
              * is not known and their order in this sequence record is
              * arbitrary. Gaps between the contigs are represented as
              * runs of N, but the exact sizes of the gaps are unknown.
              * This record will be updated with the finished sequence
              * as soon as it is available and the accession number will
              * be preserved.
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              * 1 186: contig of 186 bp in length
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              * 187 1220: contig of 1034 bp in length
              * gap of unknown length
              * 1221 2803: contig of 1583 bp in length
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              * 2804 3945: contig of 1142 bp in length
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              * 61933 67427: contig of 5495 bp in length
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              * 67428 71294: contig of 3867 bp in length
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              * 116716 132246: contig of 15531 bp in length
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              * 132247 145974: contig of 13728 bp in length
              * gap of unknown length
              * 145975 160557: contig of 14583 bp in length.
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              /clone_lib="RPC1-11 Human Male BAC"
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              Quality: 46.00 Length: 15
              Ratio: 3.538 Gaps: 0
              Percent Similarity: 86.667 Percent Identity: 60.000
              alignment_block:
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              Align seg 1/1 to reverse of: AC011694 from: 1 to: 160557
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              |||||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||
              88764 GCATATAGATCTTAATATGTTTGTGTAGACTTACGCTCAAGTAT 88720
              seq_name: gb_bai:BSUB0011
              seq_documentation_block:
              LOCUS BSUB0011 207730 bp DNA BCT 26-NOV-1997
              DEFINITION Bacillus subtilis complete genome (section 11 of 21): from 2000171
              to 2207900.
              ACCESSION 299114 AL009126
              VERSION 299114.1 GI:2634230
              KEYWORDS
              SOURCE Bacillus subtilis.
              ORGANISM Bacillus subtilis

```

Bacteria; Firmicutes; Bacillus/Clostridium group; Bacillaceae;

Bacillus.

1 (bases 1 to 207730)

Kunst, F., Ogasawara, N., Moszer, I., Albertini, A. M., Alloni, G., Azevedo, V., Bertero, M. G., Bessieres, P., Bolotin, A., Borchert, S., Boriss, R., Boursier, L., Brans, A., Braun, M., Brignell, S. C., Bron, S., Brouillet, S., Brusch, C. V., Caldwell, B., Capuano, V., Carter, N. M., Choi, S. K., Codani, J. J., Connerton, I. F., Cummings, N. J., Daniel, R. A., Denizot, F., Devine, K. M., Dusterhoft, A., Ehrlich, S. D., Emerson, P. T., Entzian, K. D., Errington, J., Fabret, C., Ferrari, E., Foulger, D., Fritz, C., Fujita, M., Fujita, Y., Fuma, S., Galizzi, A., Galleron, N., Gilm, S. Y., Glaser, P., Goffeau, A., Gollightly, E. J., Hendaut, G., Guisepi, G., Guy, B. J., Haga, K., Halech, J., Harwood, C. R., Henaut, A., Hilbert, H., Holsappel, S., Hosono, S., Hullo, M. F., Itaya, M., Jones, L., Joris, B., Karamata, D., Kasahara, Y., Klay-Bianchard, M., Klein, C., Kobayashi, Y., Koetter, P., Koningsstein, G., Krogh, S., Kumano, M., Kurita, K., Lapidus, A., Lardinois, S., Lauber, J., Lazarevic, V., Lee, S. M., Levine, A., Liu, H., Masuda, S., Maue, C., Medigue, C., Medina, N., Mellado, R. P., Mizuno, M., Moestl, D., Nakai, S., Noback, M., Noone, D., O'Reilly, M., Ogawa, K., Ogiwara, A., Oudega, B., Park, S. H., Parro, V., Pohl, T. M., Portetelle, D., Porwollik, S., Prescott, A. M., Prescan, E., Pujic, P., Purnelle, B., Rapoport, G., Rey, M., Reynolds, S., Rieger, M., Rivolta, C., Roche, B., Rose, M., Sadale, Y., Sato, T., Scanlan, E., Schleich, S., Schroeter, R., Scoffone, F., Sekiguchi, J., Sekowska, A., Seror, S. J., Serror, P., Shin, B. S., Soldo, B., Sorokin, A., Tacconi, E., Takagi, T., Takahashi, H., Takemaru, K., Takeuchi, M., Tanakoshi, A., Tanaka, T., Terpstra, P., Tognoni, A., Viari, A., Wambutt, R., Wedler, E., Wedler, H., Weitzenecker, T., Winters, P., Wipat, A., Yamamoto, H., Yamane, K., Yasumoto, K., Yata, K., Yoshida, K., Yoshikawa, H. F., Zumstein, E., Yoshikawa, H. and Danchin, A.

The complete genome sequence of the gram-positive bacterium

Bacillus subtilis

Nature 390 (6657), 249-256 (1997)

98044033

2 (bases 1 to 207730)

Kunst, F., Ogasawara, N., Yoshikawa, H. and Danchin, A.

Direct Submission

Submitted (18-NOV-1997) I. Moszer, A. Danchin, Institut Pasteur,

Regulation de l'Expression Genetique, 28 rue du Docteur Roux, 75724

Paris Cedex 15, FRANCE. E-mail: moszer@pasteur.fr,

adanchin@pasteur.fr Phone: +33 (0)1 45 68 84 41, Fax: +33 (0)1 45

68 89 48

Location/Qualifiers

1..207730

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/strain="168"

/db_xref="taxon:1423"

complement(10..1436)

/gene="yoeA"

complement(10..33)

/gene="yoeA"

complement(45..1436)

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/note="similar to hypothetical proteins from B. subtilis"

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TLFRGTGDSKTPFTLLVSTVINTIALPLVLGFPKLGISYATVISTITATE
VLWYLRKRHPLDFTVRRLKDLVLRGLVPASINNLVLSLSEIAVISEV
NLVYGSNTAAVGVNVAVSVMPSVLSIAVSTFAAQSGANEDFLKQVIRVIGWL
NYIIGVLIILIVYHQILSLFTLQESLYIAHRLMLTWSLVLPFGNAQLSATNR
ASGVLPNTVISIFAIWGVVEVPAFLVSHYTKLEILGWVGVYPAFAVSLLLIYGYQ
FWKKKQITRLIQ"
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1722..2267

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/db_xref="SPTREMBL:O34841"

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RKTYFVYKNNKWKVNOFDAVI"
2274..2292

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2319..2345

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2339..2434

/gene="trnSL-Arg1"

/product="transfer RNA-Arg"

2359..2434

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/protein_id="CAB13732.1"

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LPQKDVQNKDHLWATESONEKEKDSYFRIAEYTGNIYRERHEGKRISF"
complement(3327..3347)

terminator

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complement(3327..3578)

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complement(3348..3578)

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3763..5526

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3763..5526

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/note="alternate gene name: pac"

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/db_xref="GI:2634235"

/db_xref="SWISS-PROT:P54422"

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WGQYQGLQIATTPPPSGGIFLIQMLKILDFHNLSDQDVRSWEKYLQLAETMHLSTAD
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EOPKDYGGTQHTFVADRGNVSVYTTIEQLFGTGMVDPYGVILNNELTDFDAIP
EAGVQPNKRPLSSMTPTLFFKDDKPLVTVGSGGATIISSVLOTILYHYEGMELK
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complement(5535..6484)

gene

TMRLKENPYPVPNPIFLDTCGVOHFAAMLDLDELGLKGYVNLINSGESVNDFFYSOL
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EKIVISGKEIKVYDVLPKNGDEVVLDTFEVTLASIINDNIFNOFKLHSITDYDL
TRAKIYKLDIPDLSWSTPDGLELTORYNLSKVKKLTINELGNRTAVLRSWNEKDS
RREVQALPINIHSLSHSLTMIIDSWDSILPIHECFGHPNDMYKLAQVRECFIL
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7278. .8642
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repeat_unit

misc_feature

BASE COUNT 3042 a 1327 c 1277 g 2996 t

ORIGIN

/note="site of terminal protein"

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Quality:	45.00	Length:	12
Ratio:	3.750	Gaps:	0
Percent Similarity:	100.000	Percent Identity:	75.000

alignment_block:

US-08-653-294-21 x NIMIDRPM/rev ..

Align seg 1/1 to reverse of: NIMIDRPM from: 1 to: 8642

3 ArgLeuLeuLysValIleArgIleValLeuLys 14

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5041 AGATTGTTAATAAATGCTGAGGATTTGTTAAAA 5006

seq_name: gb_pll:SC9571X

seq_documentation_block:

LOCUS	SC9571X	29366 bp	DNA	PLN	11-AUG-1997
DEFINITION	S cerevisiae chromosome XIII cosmid 9571.				
ACCESSION	Z49810	Z71257			
VERSION	Z49810.1	GI:854472			

KEYWORDS ERG6; lactoyl glutathione lyase; MRPL39; PAR1; PDR4; PP21; protein phosphatase; ribosomal protein; SED5; SNQ3; SPT5; transfer RNA-Ser; YAP1. initiation protein; baker's yeast.

SOURCE

ORGANISM Saccharomyces cerevisiae
Eukaryota; Fungi; Ascomycota; Hemiascomycetes; Saccharomycetales;
Saccharomycetaceae; Saccharomycetes.

REFERENCE

AUTHORS 1 (bases 1 to 29366)

JOURNAL Gentles,S. and Bowman,S.

REFERENCE Unpublished

2 (bases 1 to 29366)

Barrell,B.G., Rajandream,M.A. and Walsh,S.V.

Direct Submission

Submitted (02-JUN-1995) Saccharomyces cerevisiae chromosome XIII

sequencing project, Sanger Centre, Hinxton Hall, Hinxton, Cambridge

CB10 1RQ E-mail: barrell@sanger.ac.uk

Notes:

All CDS over 100 codons have been analysed. CDS that are completely overlapped and those that are overlapped by more than 50% of their length by a larger CDS have been omitted from this analysis.

Details of the omitted CDS are available on request. The more significant matches with motifs in the PROSITE database are also included but some of these may be fortuitous. The length in codons and the calculated codon adaptation index (CAI) is given for each CDS.

Cosmid 9571 overlapped at 5' end by cosmid 8337, EMBL entry SC8337, accession no. Z46659, and at the 3' end by cosmid 8270, EMBL entry SC8270, accession no. Z48613.

FEATURES

source

1. .29366

/organism="Saccharomyces cerevisiae"

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/db_xref="taxon:4932"

/chromosome="XIII"

/clone="cosmid 9571"

/map="13L"

misc_feature

1. .463

/note="overlap with cosmid 8337, positions 34044 34506

EMBL:SC8337, Z46659"

1057. .2793

/note="YM9571.01, unknown, len: 578, CAI: 0.14"

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/product="unknown"

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YTSRDMKILNYWTTPLTKIYHITTAPEFDEKDYSTVKLLTDPKDDAGKPFITKT

QRKSNPFGSAKPDVTSQKILDEERKMENLHVEDTTTLRASLIPSSDSMATATGSKI

TILKQTPTEESHESATPTPKPLSYSEVVVERSVNNTSKGTPLSLDSPALQSKP

DKSDFKGDQGEFGKDDKAQLDYSNKDQKSETDVKQFTFKVVERHSHSRKY

NGNHNNGNFRGSRNYRGPGNGSSYKGGHNNRGNKGGYRGSSYNNNNNTNDNNN

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complement(3520. .5598)

/gene="PP21"

/db_xref="SGD:S0004478"

complement(3520. .5598)

/gene="PP21"

/note="YM9571.02c, PP21 gene, len: 692, CAI: 0.17,

SW:PP21_YEAST P26570 serine/threonine protein phosphatase

PP-21; contains PS00125 Serine/threonine specific protein

phosphatases signature"

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/db_xref="SGD:S0004478"

/product="Pp2lp"

/protein_id="CAA89936.1"

/db_xref="GI:854474"

/db_xref="SWISS-PROT:P26570"

/translation="MGNSSSKSKSDSHSSNRNRPQVSRTRTSHSVKSAKNSKS

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YSSTYENALTDNDNDKNDISHTKFRSRSSNRSPSSIRSGSVSRKSDVTHEEPNN

GSYSSNQNLYLQALTRSHASSLSHSRKSFGSGTAYSTPLNSPGLSKLTDHSG

EYFTSNTSLNHHSSRDYPSKHSINDDDIENSQLSNHSIHAEMENNDKNNTDSK

KDPNEFNDIMQSSGNKAPKFKKIDIDETIQKLDAGYAAKRTKNVCLKNNEILQ

ICIKAREIFLSPSLLEPPVKVGVGHQGDLLRFTKCGFPSSNYSYFLFGDFTD

RGQSLETILLFKYIKYENFELRGHECAVDEIRHVVRPDPDFGLINDLLMSDPTD

DFTNPLAAIVAGKIFCVHGLSPVLNSMDEIRHVVRPDPDFGLINDLLMSDPTD

SPNEDNERYGVCYKVAINKFLNKFGLDLCRAHVVVEDGYEFENDRSLVTVFSA

PNYCGEDNWGAVMSYSEGLSCFELLDPLDSALKQVKKGRKRLANQQQMMET

SITNDNESQQ"

complement(4156. .4173)

/gene="pp21"

/note="PS00125 Serine/threonine specific protein

phosphatases signature"

/db_xref="SGD:S0004478"

complement(6051. .7091)

/note="YM9571.03c, unknown, len: 346, CAI: 0.16"

/codon_start=1

/product="unknown"

/protein_id="CAA89937.1"

/db_xref="GI:854475"

/db_xref="SWISS-PROT:Q04226"

/translation="MTEPQGLDITPKVNYPPILTIANVFSTKQMDQVISEDQDYVT

WKQLNRTGTSINNLKRYKYKQRTINQODPDSINKPENLIFPODILQOQTQN

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NOTPISNIYFLQAVGKIYAGIETIELAMIVKKNLTSQMCIEFDKRTKIGYKKLYLK

KLTFSIENQYKQDYQSDSPDEDFYEDDEVDKRETTLGNLSLQSKSQSDHN

SQDKLQLQLQKLVQLQNKLQNDVSIKYNNSPLLEHREARWLRLQSDTLPNAYW

RTGEGGGSMTF"

7287. .8126

/note="YM9571.04, unknown, len: 279, CAI: 0.13, similar to

WP:C14B1.5 CE00902 Diphtheria toxin resistance protein

(39.4% identity in 155 aa overlap) and WP:C35D10.12

CE01193, (30.8% identity in 208 aa overlap)"

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EODFVPMVLKPKSKPKTKTPPAKVTKRPNLMNTPPKERSEYLRQWKEOQORSK
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QKR"
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PS00225 Crystallins beta and gamma 'Greek key' motif
signature"
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CTLLSHISGGSGRPKISKVKAFTREHIDPLAELIGDTKDKAFCEKSKAPDI
SFNALRAKEEFAMLLVGDYTDITDIDVNSKLLLEKILLNKTLOYLKIDN
DLIYLCVHELEPWLVAQLGVNTPPEIFLIANVANKASHETLPSQRLSILGKLKV
NSLNRFLQSLTNVVEKTPPELVNKTENHELMRSREIKKLOEDAYKKSLEMDRIKAIE
KESLKAQDLKLNSTARQLKWLKACIDEIQFTTGTQATLOFTSSGKRFVKKFPS
MTLLYQVSGICHYILAVYSSDPAENSNALQDKIRQLSADDDMLCFEGOLETATAT
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LVEALDEEEDENEEO"
10178..10813
/note="YM9571.06, len: 211, CAI: 0.22, potential
transmembrane protein, similar at C-terminus to
SW:G25L_CANFA P27869 glycoprotein 25L precursor (24.4%
identity in 168 aa overlap) and to SW:P24B_YEAST P32803
P24B protein precursor (26.0% identity in 131 aa overlap)"
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QAQYRGSLRAIELDTESGAEDWKNKPIEVLRRVEETIDIVDELTY
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FLPSVATQITWGVRCRPGKEKELIRKLKKFNLDKRAMKKKILKLSIFORDNVTGR
IYIEAPQSVIEKFCNGVPDIYSQKLLIPVQELPDLKPNKSDVALEESYVRIKR
GIYKGLDAMWDOIENNLEVKIPLRDIYQKDEIDPTTQOKSRRTFAHRAPPQL
FNPMAURLDQANLYKRDHFTYKNEYIDGYLYKSFRIQHVETKNTQPTVEELARE
KSGKGVADLTVSQSIRKAAKAVTFQPDRIEVLNGEQRSKGTIVTTRTDIATIKL
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ITNNLSKIDTTATSEYALDIVELSAKNVACIIQAGHDIKVIDETGKVTITKG
SILSKINTARARVSVDPANGNEIKGDTIVEKYSRREGQVLYIQTOQIFVYVKKIIVE
NAGVFVNPFSVEAVASNDKSNKMDLSKMPPELISGPPSPSKTFPOQIQSRGREGI
VALGKTVIRISAGYKQGLGVKDVNGDKATVELHSKNKHITIDKHLIYINREGGEGI
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  Ratio: 3.462       Gaps: 0
  Percent Similarity: 92.857   Percent Identity: 57.143

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seq_name: gb_hgt5.AC015353

seq_documentation_block:
  LOCUS AC015353 29459 bp DNA HTG 16-NOV-1999
  DEFINITION Drosophila melanogaster, *** SEQUENCING IN PROGRESS ***, in ordered
  pieces.
  ACCESSION AC015353
  VERSION AC015353.1 GI:6435982
  KEYWORDS HTG; HTGS_PHASE2.
  SOURCE fruit fly.
  ORGANISM Drosophila melanogaster
    Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
    Pterygota; Neoptera; Endopterygota; Diptera; Brachycera;
    Muscomorpha; Ephydroidea; Drosophilidae; Drosophilla.
  REFERENCE 1 (bases 1 to 29459)
  AUTHORS Adams,M. and Venter,J.C.
  TITLE Direct Submision
  JOURNAL Submitted (16-NOV-1999) Celera Genomics, 45 West Gude Drive,
  Rockville, MD, USA
  COMMENT This sequence was identified as CDM:10210956 by the submitter.
  For further information on this sequence e-mail to fly@celera.com.
  * NOTE: This is a 'working draft' sequence.
  * This sequence will be replaced
  * by the finished sequence as soon as it is available and
  * the accession number will be preserved.
  FEATURES
    source
      location/Qualifiers
        1..29459
        /organism="Drosophila melanogaster"
        /db_xref="taxon:7227"
  BASE COUNT 7846 a 6592 c 6879 g 8142 t
  ORIGIN

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  Ratio: 3.750       Gaps: 0
  Percent Similarity: 100.000   Percent Identity: 58.333

alignment_block:
  US-08-653-294-21 x AC015353/rev ..
  Align seg 1/1 to reverse of: AC015353 from: 1 to: 29459
  1 AlaTyrArgLeuLeuLysValIleArgIleVal 12
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1853 GCCTACCGCTCGTACGATCGTTCGGATCGTC 1818

seq_name: gb_pl1.SPAC9G1

seq_documentation_block:

LOCUS SPAC9G1 30985 bp DNA PLN 09-SEP-1998
 DEFINITION S.pombe chromosome I cosmid c9G1.
 ACCESSION 298763
 VERSION 298763.1 GI:2370549

KEYWORDS beta-transducin; cdc12 homolog; cytochrome oxidase biogenesis protein; Homol D box; inositol metabolism; Inositol polyphosphate phosphatase; lysophospholipase; MAP kinase kinase kinase; oxal; ribosomal protein L30e; rpl30; septin homolog; serine threonine protein kinase; spn4; src homology domain; vacuolar sorting; waki; WD repeat.

SOURCE fission yeast.

ORGANISM

Schizosaccharomyces pombe
 Eukaryota; Fungi; Ascomycota; Archiascomycetes;
 Schizosaccharomycetales; Schizosaccharomycetaceae;
 Schizosaccharomycetes.

REFERENCE

AUTHORS Churcher,C.M. and Gentles,S.

JOURNAL Unpublished

2 (bases 1 to 30985)

Barrell,B.G., Rajandream,M.A. and Wood,V.

AUTHORS

Submitted (26-AUG-1997) Schizosaccharomycetes pombe chromosome I
 sequencing project, Sanger Centre, Wellcome Trust Genome Campus,
 Hinxton, Cambridge CB10 1SA E-mail: barrell@sanger.ac.uk

COMMENT

Details of yeast sequencing at the Sanger Centre are available on
 the World Wide Web.

(URL: <http://www.sanger.ac.uk/Projects/S.pombe/>)
 Protein coding regions (CDS) have been predicted with the help of
 computer analysis using the Genefinder program in FomBase (an ACEDB
 database) with additional predictions for the branch-acceptor sites
 supplied by the program Sp3splice. CAUTION: It is possible that for
 any individual CDS we may have underestimated or overestimated the
 number of introns/exons or we may not have chosen the correct
 splice donor/acceptor sites. CDS are numbered using the following
 system eg SPAC9G1.01c. Sp (S. pombe), A (chromosome 1), c5H10
 (cosmid name), .01 (first CDS), c (complementary strand).
 The more significant matches with motifs in the PROSITE database
 are also included but some of these may be fortuitous. The length
 in codons is given for each CDS.

IMPORTANT: This sequence MAY NOT be the entire insert of the
 sequenced clone. It may be shorter because we only sequence
 overlapping sections once, or longer, because we arrange for a
 small overlap between neighbouring submissions. Cosmid c9G1
 overlaps cosmid c3H1 at its 3' end.

FEATURES

source

Location/Qualifiers

1..30985

/organism="Schizosaccharomycetes pombe"

/strain="972h"

/db_xref="taxon:4896"

/chromosome="I"

/clone="cosmid c9G1"

/map="I"

1..476

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1..461

misc_feature

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/note="nominal overlap with EM_FUN:SPAC3H1 Z68144
 position:36460..36920"

<1..476

CDS

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 /note="SPAC9G1.01, similarity: to YGR066C, YG29_YEAST,
 P53242, hypothetical 34.0 kd protein, (292aa), fasta
 scores, opt:210, E():4e-09, (32.7% identity in 159 aa
 overlap) similarity: to YBR105C, YBV5_YEAST, P38263,
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 AQAEMKLPYEDKLLPLVPFIKLEALRLAYAKKIKGNPLIIVLMDLDFKI
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 SITTVYKEFIRLSKLCMRISSTVDCVSAREACGVNCHDLIVHVFSAAEFGRI
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 NILLDDANFLAIRKVGKSMALITHEDVLGAKSKVAALKQRESEVSSPRLTSFG
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 VLERLNHPNVYTYGVEVHREKVIYFMFCQSGSLADLHAGRIEDENLVKVVYVOLL
 EGLAYTHSHOILHRDIPANILIDHRGMIKYSGFSGALYVSPTEDEYVEDIQPELQ
 HLAGPTMYMAPEIILGKKGDFGADWSLGCVILEMTGTSPWSEMDNEWALMYHVA
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 /db_xref="SWISS-PROT:P52808"
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 complement(6772..6846)

misc_feature

misc_feature

gene

CDS

misc_feature

misc_feature

misc_feature

misc_feature

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AIFLPTQSAVEFFSYAIRKMSRLSDVGTFTGGLAWFKDLSIPDPYILPINAGL
MFSGMNMRANTASTIGNTNWRTFFFLCCLLSPLLTAKLPAAIFMWIPSSLFNIQV
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9054..9072
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9769..11928
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10423..10468,10518..11459,11506..11928)
/genes="SPAC9G1.05"
/notes="SPAC9G1.05, len:595aa, similarity: to WD42_DICDI,
P54686, wd40 repeat protein 2, (597aa), fasta scores,
opt:1374, E():0, (36.7% identity in 603 aa overlap), also
similar eg. YMR092C, AIP1_YEAST, P46680, actin interacting
protein 1, (615aa), fasta scores, opt:94, E():0, fasta
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overlap)"
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KTGDQVEIDAHGSGIFSISWSDSOFVTSSAGYSCKIWDANTGLIREWLSDDKKQ
LVCTVPTKDLIIIVNSKGLNLTYNPDKCKVIDTYIGHORSITAAATLSPDCHFYTAS
YDGTVLISWDIGKKAFLPGESTNQMNMADDDHVTIGMDTLRLVIDIKQCCFAK
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alignment_scores:
  Quality: 45.00      Length: 14
  Ratio: 3.750      Gaps: 0
  Percent Similarity: 85.714      Percent Identity: 50.000
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alignment_block:
  US-08-653-294-21 x SPAC9G1
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Align seg 1/1 to: SPAC9G1 from: 1 to: 30985
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12984 TATAAACTGCTGGTCCGCGATTCTCAGGTACGTAGTAAATAT 13025
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seq_name: gb_inl:CELF2969
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seq_documentation_block:
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LOCUS      CELF2969      42751 bp      DNA      INV      07-AUG-1997
DEFINITION Caenorhabditis elegans cosmid F2969.
ACCESSION  AF016440
VERSION    AF016440.1  GI:2315352
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```
KEYWORDS
SOURCE
ORGANISM
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Caenorhabditis elegans strain=Bristol N2.
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Caenorhabditis elegans
Eukaryota; Metazoa; Nematoda; Secernentea; Rhabditia; Rhabditida;
Rhabditina; Rhabditoidea; Rhabditidae; Peloderinae; Caenorhabditis.
1 (bases 1 to 42751)
```

```
AUTHORS
```

```
Wilson,R., Ainscough,R., Anderson,K., Baynes,C., Berks,M.,
Bonfield,J., Burton,J., Connell,M., Copsey,T., Cooper,J.,
Coulson,A., Craxton,M., Dear,S., Du,Z., Durbin,R., Favello,A.,
Johnston,L., Jones,M., Kershaw,J., Kirsten,J., Laister,N.,
Latreille,P., Lightning,J., Lloyd,C., McMurray,A., Mortimore,B.,
O'Callaghan,M., Parsons,J., Percy,C., Rifken,L., Roopra,A.,
Saunders,D., Showkneen,R., Smaldon,N., Smith,A., Sonnenhammer,E.,
Staden,K., Sulston,J., Thierry-Mieg,J., Thomas,K., Vaudin,M.,
Vaughan,R., Waterston,R., Watson,A., Weinstock,L.,
Wilkinson-Sproat,J. and Wohldman,P.
2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
elegans
Nature 368 (6466), 32-38 (1994)
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TITLE
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JOURNAL      Nature 368 (6466), 32-38 (1994)
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MEDLINE      94150718
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REFERENCE    2 (bases 1 to 42751)
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```
AUTHORS      Langston,Y.
```

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TITLE        The sequence of C. elegans cosmid F2969
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```
JOURNAL      Unpublished (1997)
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```
REFERENCE    3 (bases 1 to 42751)
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```
AUTHORS      Waterston,R.
```

```
TITLE        Direct Submission
```

```
JOURNAL      Submitted (30-JUL-1997)
```

```
COMMENT      Submitted by:
```

```
Genome Sequencing Center
```

```
Department of Genetics, Washington University,
```

```
St. Louis, MO 63110, USA, and
```

```
Sanger Centre, Hinxton Hall
```

```
Cambridge CB10 1RQ, England
```

```
e-mail: rwenematode.wustl.edu and jes@sanger.ac.uk
```

NOTICE: This sequence may not be the entire insert of this clone. It may be shorter because we only sequence overlapping sections once, or longer because we provide a small overlap between neighboring submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded or sequenced with an alternate chemistry; an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone

NEIGHBORING COSMID INFORMATION:

The 5' cosmid is C10F3, 200 bp overlap; 3' cosmid is R01B10, 2200 bp overlap. Actual start of this cosmid is at base position 197 of CELF2969; actual end is at 40555 of CELF2969

NOTES:

Coding sequences below are predicted from computer analysis, using the program Genefinder(P. Green and L. Hillier, ms in preparation).

Location/Qualifiers

1. .42751

/organism="Caenorhabditis elegans"

/strain="Bristol N2"

OM of: US-08-653-294-21 to: N_Geneseq_36:* out_format : pfs

Date: Feb 8, 2000 7:30 PM

About: Results were produced by the GenCore software, version 4.5,
Copyright (c) 1993-2000 Compugen Ltd.

Command line parameters:

```
-MODEL=frame p2n.model -DEV=xlp  
-O=/cnp1.1/USPTO.SPOOL/US08653294/runat_04022000.160701.15807/app_query.fasta.2  
-DB=N_Geneseq_36 -QFMT=fastap -SUFFIX=ring -GAPOP=12.000  
-GAPEXT=4.000 -MINMATCH=0.100 -LOOPCL=0.000 -LOOPEXT=0.000  
-QGAPOP=4.500 -QGAPEXT=0.050 -XGAPOP=10.000 -XGAPEXT=0.500  
-XGAPEXT=6.000 -XGAPEXT=7.000 -XGAPEXT=10.000 -XGAPEXT=0.500  
-DELOP=6.000 -DELEXT=7.000 -START=1 -MATRIX=biosum62  
-TRANS=human40.cdi -LIST=45 -DOCALIGN=200 -THR SCORE=pct  
-ALIGN=15 -MODE=LOCAL -OUTFMT=pfs -NORM=ext -MINLEN=0  
-MAXLEN=100000 -USER=US08653294 -NCPU=6 -ICPU=3 -NO_XLPXY -WAIT  
-THREADS=1
```

Search information block:

Query: US-08-653-294-21

Query length: 15

Database: N_Geneseq_36:*

Database sequences: 311585

Database length: 125096042

Search time (sec): 873.190000

score_list:

Sequence	Strd Orig	zscore	Escore	Len	Documentation
N_Geneseq_36:X13927	+	42.00	125.48	21.76	810 ! H. pylori GHPO 635 gene. New is
N_Geneseq_36:O56758	+	42.00	116.44	69.39	2255 ! Chitinase derivative gene #1
N_Geneseq_36:X13348	+	41.00	108.74	186.35	3721 ! Enterococcus faecalis genome d
N_Geneseq_36:T33641	+	40.00	118.50	54.67	869 ! Aspergillus arabinofuranosidas
N_Geneseq_36:T33646	+	40.00	108.78	185.41	2555 ! Aspergillus arabinofuranosidas
N_Geneseq_36:X13195	+	40.00	97.67	770.82	8991 ! Enterococcus faecalis genome d
N_Geneseq_36:X30892	+	39.00	118.40	54.01	593 ! Streptococcus pneumoniae poly
N_Geneseq_36:V42962	+	39.00	107.77	210.96	1975 ! Streptococcus pneumoniae poly
N_Geneseq_36:V07122	+	39.00	105.34	288.02	2600 ! Cytophaga drobachiensis kappa
N_Geneseq_36:T84234	+	39.00	103.76	352.79	3110 ! DNA encoding an autolysin and
N_Geneseq_36:V53357	+	39.00	103.76	352.79	3110 ! DNA encoding 2 Staphylococcus
N_Geneseq_36:V52309	+	39.00	101.58	467.01	3984 ! Streptococcus pneumoniae genom
N_Geneseq_36:V59717	+	39.00	101.47	473.26	4031 ! Tumour rejection antigen precu
N_Geneseq_36:V69720	+	39.00	101.06	499.13	4225 ! Tumour rejection antigen precu
N_Geneseq_36:V74331	+	39.00	93.04	1.4e+03	10470 ! Staphylococcus aureus contig
N_Geneseq_36:V70401	+	39.00	89.60	2.2e+03	15462 ! Human parainfluenza virus typ
N_Geneseq_36:X20248_04	+	39.00	72.28	2.0e+04	110000 ! Continuation (5 of 10) of
N_Geneseq_36:X20248_08	+	39.00	72.28	2.0e+04	110000 ! Continuation (5 of 10) of
N_Geneseq_36:X20248_08	+	39.00	72.28	2.0e+04	110000 ! Continuation (9 of 10) of
N_Geneseq_36:Q03147	+	38.00	105.55	280.35	1751 ! Yeast Ubiquitin hydrolase gene
N_Geneseq_36:X02969	+	38.00	105.40	285.79	1781 ! Human IL-1ra BAC contiguous DN
N_Geneseq_36:X03047	+	38.00	90.95	1.8e+03	9158 ! Human IL-1ra BAC contiguous DN
N_Geneseq_36:V74364	+	38.00	85.70	3.6e+03	16592 ! Staphylococcus aureus contig
N_Geneseq_36:V52273	+	38.00	80.80	6.7e+03	28882 ! Streptococcus pneumoniae gen
N_Geneseq_36:T58840_4	+	38.00	69.00	3.0e+04	110000 ! Continuation (5 of 6) of T5
N_Geneseq_36:V21209_10	+	38.00	69.00	3.0e+04	110000 ! Continuation (11 of 17) of
N_Geneseq_36:X20248_06	+	38.00	69.00	3.0e+04	110000 ! Continuation (7 of 10) of
N_Geneseq_36:X20250	+	38.00	68.89	3.0e+04	111309 ! Borrelia burgdorferi polynucl
N_Geneseq_36:V74525	+	37.50	90.76	1.9e+03	7769 ! Staphylococcus aureus contig S
N_Geneseq_36:X00666	+	37.00	107.12	229.22	1011 ! Human secreted protein gene 56
N_Geneseq_36:V69759	+	37.00	106.78	239.52	1051 ! Arabidopsis ovary-specific AGU
N_Geneseq_36:V29580	+	37.00	105.71	274.65	1186 ! Pythium oligandrum isolate 23-
N_Geneseq_36:X06780	+	37.00	102.29	426.14	1748 ! Human testis secreted protein
N_Geneseq_36:X13220	+	37.00	96.20	930.92	3485 ! Enterococcus faecalis genome d
N_Geneseq_36:X13280	+	37.00	95.19	1.1e+03	3907 ! Enterococcus faecalis genome d
N_Geneseq_36:V52207	+	37.00	82.45	5.4e+03	16535 ! Streptococcus pneumoniae gen
N_Geneseq_36:X13336	+	37.00	76.41	1.2e+04	32768 ! Enterococcus faecalis genome
N_Geneseq_36:V30458_3	+	37.00	65.72	4.5e+04	110000 ! Continuation (4 of 6) of V3
N_Geneseq_36:V21209_01	-	37.00	65.72	4.5e+04	110000 ! Continuation (2 of 17) of
N_Geneseq_36:X20248_08	-	37.00	65.72	4.5e+04	110000 ! Continuation (9 of 10) of
N_Geneseq_36:V30459_3	-	37.00	65.72	4.5e+04	110000 ! Continuation (4 of 6) of V3

N_Geneseq_36:Q46292 - 36.50 101.95 444.90 1508 ! PAL regulator gene. Regulat
N_Geneseq_36:Q46293 - 36.50 92.47 1.5e+03 4415 ! Phenylalanine ammonia lyase
N_Geneseq_36:T78725 - 36.00 127.42 16.97 70 ! SELEX generated ligand to hCG
N_Geneseq_36:V89042 + 36.00 114.00 94.88 320 ! EST clone BV31. New polynuc

seq_name: N_Geneseq_36:X13927

seq_documentation_block:

```
ID X13927 standard; DNA; 810 BP.  
AC X13927;  
CT 31-MAR-1999 (first entry)  
DE H. pylori GHPO 635 gene.  
KW GHPO protein; Helicobacter infection; gastroduodenal disease; gastritis;  
KW peptic ulcer disease; ss.  
FH Helicobacter pylori.  
FT key Location/Qualifiers  
FT CDS 98..760  
FT /tag= a
```

WO9843478-A1.

PD 08-OCT-1998.

PF 01-APR-1998; U06371.

PR 29-JUL-1997; US-902615.

PR 01-APR-1997; US-833457.

PR 24-JUN-1997; US-881227.

PA (HUMA-) HUMAN GENOME SCI INC.

PA (INMR) MERIEUX ORAVAX PASTEUR MERIEUX SERUMS.

PI Al-Garawi A, Kleanthous H, Miller C, Oomen RP, Tomb J;

DR WPI: 98-542293/46.

DR P-PSDB: W98208.

PT New isolated Helicobacter polynucleotides - used to develop products

PT for the diagnosis, prevention and treatment of Helicobacter

PT infections and gastrointestinal diseases

PS Claim 1: Page 111-113; 2054pp; English.

CC This sequence represents a polynucleotide of the invention. It was

CC isolated from Helicobacter pylori and encodes a H.pylori GHPO protein.

CC The polypeptides can be used for preventing or treating Helicobacter

CC infections, and gastroduodenal diseases associated with these

CC infections, including acute, chronic, and atrophic gastritis, and peptic

CC ulcer diseases, e.g. gastric and duodenal ulcers. They can also be used

CC for the production of antibodies. The products can also be used for

CC detection and diagnosis.

CC Sequence 810 BP; 223 A; 123 C; 188 G; 276 T;

alignment_scores:

Quality: 42.00 Length: 15

Ratio: 3.231 Gaps: 0

Percent Similarity: 86.667 Percent Identity: 53.333

alignment_block:

US-08-653-294-21 x X13927 ..

Align seg 1/1 to: X13927 from: 1 to: 810

```
1 AlaTyrArgLeuLeuLysValleLysValleArgLleValleLysTyr 15  
||||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
720 GCTTACCGGTAGCTATTTCATATATATATATATATATATATATATAT 764
```

seq_name: N_Geneseq_36:Q56758

seq_documentation_block:

ID Q56758 standard; DNA; 2255 BP.

AC Q56758;

CT 12-OCT-1994 (first entry)

DE Chitinase derivative gene #1.

KW Chitinase; derivative; beta-1,4-glycoside bond; chitin;

KW beta-N-acetylhexosaminidase activity; yeast; ss.

OS Rhizopus niveus.

FH key Location/Qualifiers

FT cds 138..1750

FT /tag= a

FT /product= Chitinase

FT exon 138..242

FT intron /*tag= b
 FT 243..305
 FT /*tag= c
 FT exon 306..498
 FT /*tag= d
 FT intron 499..566
 FT /*tag= e
 FT exon 567..1750
 FT /*tag= f

PN J06046849-A.
 PD 22-FEB-1994.
 PF 28-JUL-1992; 201427.
 PR 28-JUL-1992; JP-201427.
 PA (KAGO) KAGOME KK.
 DR WPI; 94-097015/12.
 DR P-FSDB; R48669.
 PT New chitinase enzyme and coding sequence - cleaves beta-1,4-glycoside
 PT bond of chitin but has no beta-N-acetylhexosaminidase activity
 PS Claim 11; Page 21-23; 66pp; Japanese.
 CC The sequences given in Q56756-61 encode chitinase derivatives which
 CC cleave the beta-1,4-glycoside bond of chitin but have substantially
 CC no beta-N-acetylhexosaminidase activity. These sequences may be
 CC introduced into yeast and cultured for the production of the
 CC chitinase proteins.
 SQ Sequence 2255 BP; 614 A; 448 C; 446 G; 747 T;

alignment_scores:
 Quality: 42.00 Length: 9
 Ratio: 4.667 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 100.000

alignment_block:

US-08-653-294-21 x Q56758

Align seg 1/1 to: Q56758 from: 1 to: 2255

7 LysValIleArgIleValLeuLysTyr 15

|||||
 1867 AAGTAATAGATTGCTTAAATAT 1893

seq_name: N_Geneseq_36:X13348

seq_documentation_block:

ID X13348 standard; DNA; 3721 BP.

AC X13348;

DT 19-MAR-1999 (first entry)

DE Enterococcus faecalis genome contig SEQ ID NO:411.

KW Enterococcus faecalis; contig; detection; Enterococcal infection;

KW vaccine; attenuation; computer readable medium; ds.

OS Enterococcus faecalis.

PN W0850555-A2.

PD 12-NOV-1998.

PF 04-MAY-1998; U08985.

PR 14-NOV-1997; US-066009.

PR 06-MAY-1997; US-044031.

PR 16-MAY-1997; US-046655.

PA (HUMA-) HUMAN GENOME SCI INC.

PI Barash SC, Dillon PJ, Kunsch CA;

DR WPI; 99-045171/04.

PT New isolated Enterococcus faecalis polynucleotides and polypeptides

PT - used to develop products for the detection of Enterococcus and for

PT use in vaccines for prevention or attenuation of Enterococcus

PT infection

PS Claim 1; Page 1626-1628; 2084pp; English.

CC A computer readable medium has been developed which has recorded on it

CC 982 nucleotide sequences isolated from the Enterococcus faecalis genome.

CC X12938 to X13919 represent these nucleotide sequences which are primary

CC nucleotide sequences, also known as contigs. The computer-based system

CC can identify fragments of the Enterococcus faecalis genome with

CC commercial importance. The products can be used to detect the presence

CC of Enterococcus faecalis in samples. They can also be used for

CC diagnosing Enterococcal infection in an animal and monitoring

CC progression of disease, and for identifying agents which can be used to
 CC modulate the growth or pathogenicity of Enterococcus faecalis, or
 CC another related organism, in vivo or in vitro. In particular the
 CC polypeptides encoded by the Enterococcus faecalis nucleotide sequences
 CC can be used in vaccines to prevent or attenuate an Enterococcal
 CC infection.
 SQ Sequence 3721 BP; 1387 A; 582 C; 785 G; 934 T;

alignment_scores:

Quality: 41.00 Length: 14
 Ratio: 3.417 Gaps: 0
 Percent Similarity: 85.714 Percent Identity: 50.000

alignment_block:

US-08-653-294-21 x X13348/rev

Align seg 1/1 to reverse of: X13348 from: 1 to: 3721

2 TyrArgLeuLeuLysValIleArgIleValLeuLysTyr 15

||||| : : : : : ||| : : : : : ||| : : : : : |||

698 TACCGATCCATGCTTAAATGATCAGAAATATTTCGGATAC 657

seq_name: N_Geneseq_36:T33641

seq_documentation_block:

ID T33641 standard; DNA; 869 BP.

AC T33641;

DT 11-DEC-1996 (first entry)

DE Aspergillus arabinofuranosidase gene promoter.

KW Promoter; arabinofuranosidase; AbfC; arabinoxylan;

KW viscosity modifier; food; feedstuff; ss.

OS Aspergillus niger strain 3M43.

PN W09628416-A1.

PD 26-SEP-1996.

PF 11-MAR-1996; E01009.

PR 17-MAR-1995; GB-003479.

PA (DANI-) DANISCO AS.

PI Baruch A, Madrid SM, Rasmussen P;

DR WPI; 96-443191/44.

PT Aspergillus arabinofuranosidase - useful for degradation of

PT arabinoxylan

PS Claim 8; Page 48; 105pp; English.

CC The promoter (T33641) of the arabinofuranosidase gene of Aspergillus

CC niger 3M43 was identified in a full-length gene (T33646) isolated

CC from a genomic DNA library. The promoter is strongly repressed by

CC glucose and induced by intermediates of xylose metabolism. It

CC can be used to control the expression of a gene of interest, e.g.

CC A. niger arabinofuranosidase or a heterologous gene, in a host

CC cell, pref. Aspergillus or a transgenic plant.

SQ Sequence 869 BP; 224 A; 209 C; 193 G; 243 T;

alignment_scores:

Quality: 40.00 Length: 12
 Ratio: 3.333 Gaps: 0
 Percent Similarity: 100.000 Percent Identity: 58.333

alignment_block:

US-08-653-294-21 x T33641

Align seg 1/1 to: T33641 from: 1 to: 869

2 TyrArgLeuLeuLysValIleArgIleValLeu 13

||||| : : : : : ||| : : : : : ||| : : : : : |||

165 TACCAACTTTTGTGATTCGATTCAGAAATACTTTC 200

seq_name: N_Geneseq_36:T33646

seq_documentation_block:

ID T33646 standard; DNA; 2555 BP.

AC T33646;

DT 11-DEC-1996 (first entry)

DR	WPI: 99-045171/04.
PT	NT New isolated Enterococcus faecalis polynucleotides and polypeptides
PT	NT - used to develop products for the detection of Enterococcus and for
PT	use in vaccines for prevention or attenuation of Enterococcus
PT	infection.
PS	Claim 1: Page 1239-1244; 2084app; English.
CC	A computer readable medium has been developed which has recorded on it
CC	992 nucleotide sequences isolated from the Enterococcus faecalis genome
CC	X12998 to X13919 represent these nucleotide sequences which are primary
CC	nucleotide sequences, also known as contigs. The computer-based system
CC	can identify fragments of the Enterococcus faecalis genome with
CC	commercial importance. The products can be used to detect the presence
CC	of Enterococcus faecalis in samples. They can also be used for
CC	diagnosing Enterococcal infection in an animal and monitoring
CC	progression of disease, and for identifying agents which can be used to
CC	modulate the growth or pathogenicity of Enterococcus faecalis, or
CC	another related organism, in vivo or in vitro. In particular the
CC	polypeptides encoded by the Enterococcus faecalis nucleotide sequences
CC	can be used in vaccines to prevent or attenuate an Enterococcal
CC	infection.
CC	sequence 8991 BP: 2910 A: 1478 C: 1913 G: 2683 T:
SQ	

```
alignment_scores:
  Quality: 40.00      Length: 13
  Ratio: 3.333       Gaps: 0
  Percent similarity: 92.308  Percent Identity: 53.846

alignment_block:
  US-08-653-294-21 x X13195/rev ..
```

Align seg 1/1 to reverse of: X13195 from: 1 to: 8991

2 TyrArgLeuLeuLeuLysValIleArgIleValLeuLys 14
:: |||||:::|||||:::
1946 TTCGCCCTCCTCATCAAAATAGTACAGATAGTCTCGCGC 1908

seq_name: N_Geneseq_36:X30892

seq_documentation_block:
ID: X30892 standard: DNA: 593 bp.

AC	X30892;
DE	20-MAY-1999 (first entry)
DE	<i>Streptococcus pneumoniae</i> genomic DNA sequence SEQ ID NO:169.
KW	<i>Streptococcus pneumoniae</i> strain 0100993; vaccine; immune response;
KW	<i>Streptococcus pneumoniae</i> strain 0100993; vaccine; immune response;
OS	streptococcal infection; pneumococcal; ss.
OS	<i>Streptococcus pneumoniae</i> .

Streptococcal infection;
Streptococcus pneumoniae.
WO9737026-A1.

PD 09-OCT-1997.

PF 01-APR-1997; U05306.

PR 22-AUG-1996; US-025788.
 PR 03-APR-1996; US-014690

PR 02-APR-1996; US-014690.
PA (SMIK) SMITHKLINE BEECHAM CORP.

PA (SMIK) SMITHKLINE BEECHAM PLC.

PI Black MT, H

PI Stodola RK;
EXP. 07-503111AC

DR WP1; 97-503111/
DR P-PSDB: V11310

PT Nucleic acids encoding pneumococcal polypeptide(s) - useful in

PT vaccines, drug

PS Claim 5; Page 156; 354pp; English.

CC X30724 to X30946 represent genomic
CC streptococcus pneumoniae strain 01

CC streptococcus pneumoniae strain 01
CC encode the novel proteins given in

CC isolated from Streptococcus pneumoniae

CC streptococcal infections and in as

inhibit or activate the activity of

be used to treat an individual having a protein vector expressing the protein.

CC protein: Vectors expressing the pr
CC protective immune response in mamm
CC

Sequence 593 BP; 206 A; 76

This nucleotide sequence comprises novel human tumour rejection antigen precursor (TRAP) MAGE-C1 cDNA (see also V69720). MAGE-C1 is a novel member of the MAGE family that may be recognised by cytotoxic T cells, leading to lysis of the tumour cells which express it. MAGE-C1 and MAGE-C2 (see W81546-47) are expressed in variety of tumours and in normal testis cells, but not by other normal cells. The MAGE-C1 cDNA was isolated from a melanoma LB373-PCR cDNA library using a probe generated from LB-373-MEL. RNA by PCR (see V69728-29). It shows homology to MAGE-A1 cDNA (see

```
CC comprising at least one tumour rejection antigen derived from
CC MAGE-C1 or MAGE-C2 and at least one other tumour rejection antigen.
CC MAGE-C1 and MAGE-C2 can be used in a method for determining the
CC presence of cytolytic T cells specific for complexes of a human
CC leukocyte antigen (HLA).
SQ Sequence 4225 BP; 871 A; 1198 C; 923 G; 1233 T;

alignment_scores:
  Quality: 39.00      Length: 13
  Ratio: 3.000       Gaps: 0
Percent Similarity: 100.000 Percent Identity: 38.462

alignment_block:
US-08-653-294-21 x V69720 ..
Align seg 1/1 to: V69720 from: 1 to: 4225

      2 TyrArgLeuLeuIleLysValIleArgIleValLeuLys 14
      TACAGAGTTCTCTGAGTGTCTCAAGTGTCTTTGAGG 1565
seq_name: N_Geneseq_36:V74331

seq_documentation_block:
ID V74331 standard; DNA; 10470 BP.
AC V74331;
DT 16-MAR-1999 (first entry)
DE Staphylococcus aureus contig SEQ ID #20.
KW Computer readable medium; vaccine; S.aureus infection; Immunodetection;
KW cellulitis; eyelid infection; food poisoning; osteomyelitis; therapy;
KW skin infection; surgical wound infection; scalded skin syndrome;
KW toxic shock syndrome; ds.
OS Staphylococcus aureus.
FH Key Location/Qualifiers
FT misc_feature 361..420
FT /*tag=a
FT /note="these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"
FT misc_feature 2161..2220
FT /*tag=b
FT /note="these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"
FT misc_feature 3961..4020
FT /*tag=c
FT /note="these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"
FT misc_feature 5761..5820
FT /*tag=d
FT /note="these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"
FT misc_feature 7561..7620
FT /*tag=e
FT /note="these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"
FT misc_feature 9361..9420
FT /*tag=f
FT /note="these bases represent a line of missing text in
FT the sequence listing in the specification. They
FT are included to maintain the nucleotide numbering
FT given in the specification for this DNA sequence"
FN EP-786519-A2.
PD 30-JUL-1997.
```

```
PF 07-JAN-1997; 100117.
PR 05-JAN-1996; US-009861.
PA (HUMA-) HUMAN GENOME SCI INC.
PI Barash SC, Choi GH, Dillon PJ, Fannon MR, Kunsch CA,
PI Rosen CA;
DR WPI; 97-374922/35.
PT Polynucleotide(s) and proteins derived from Staphylococcus aureus
PT stored on computer readable medium and used in the production of
PT anti-S.aureus vaccines
PS Claim 1; Page 271-277; 3271pp; English.
CC This sequence represents one of 5191 Staphylococcus aureus DNA sequences
CC of the invention. The DNA sequences are recorded on a computer readable
CC medium, preferably selected from a floppy or hard disk, random access
CC memory (RAM), read-only memory (ROM) or CD-ROM. Homology searches using
CC the S.aureus DNA sequences allows putative functions to be assigned so
CC that protein-encoding or regulatory regions of commercial, therapeutic or
CC industrial importance can be obtained. Specifically, sequences which are
CC likely to encode antigens have been identified and these polypeptides can
CC be used in a vaccine composition against S.aureus infection. The
CC polypeptides can also be used in a kit for the immunodetection of
CC S.aureus in a sample. S.aureus is implicated in numerous human diseases,
CC including cellulitis, eyelid infections, food poisoning, osteomyelitis,
CC skin and surgical wound infections, scalded skin syndrome, toxic shock
CC syndrome, etc. Organisms transformed with the DNA sequences can be used
CC for recombinant production of the polypeptides. The new DNA sequences
CC (and their fragments) are useful as primers or probes for isolating
CC homologues of any of the S.aureus DNA sequences contained on the
CC computer readable medium.
SQ Sequence 10470 BP; 2804 A; 2009 C; 1550 G; 3742 T;
```

```
alignment_scores:
  Quality: 39.00      Length: 14
  Ratio: 3.250       Gaps: 0
Percent Similarity: 85.714 Percent Identity: 35.714

alignment_block:
US-08-653-294-21 x V74331/rev ..
Align seg 1/1 to reverse of: V74331 from: 1 to: 10470

      2 TyrArgLeuLeuIleLysValIleArgIleValLeuLysTyr 15
      TACCAGTAGCTTCAAGATTATGAGGTTATTTATCGGTAT 9246
```

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